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## (57) Abstract

Arays of polynucleotide spots and kits comprising the same, as well as methods for their preparation and use are provided. The subject arrays include a plurality of polynucleotide spots stably associated with the surface of a solid support. At least a portion of the polynucleotide spots comprises a polynucleotide probe composition that is made up of unique polynucleotides, where all of the unique polynucleotides of the array correspond to a common type of gene. Also provided are sets of a representational number of gene specific primers suitable for use in generating target nucleic acid for use with the subject arrays. The subject arrays find use in hybridization assays, particularly in assays for the identification of differential gene expression pattems among two or more different types of cells.

## NUCLEIC ACID ARRAYS

## CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of US Patent $5,994,076$, the disclosures of which are herein incorporated by reference.

## INTRODUCTION

## Technical Field

The field of this invention is biopolymeric arrays.

## Background of the Invention

"Biochips" or arrays of binding agents, such as oligonucleotides and peptides, have become an increasingly important tool in the biotechnology industry and related fields. These binding agent arrays, in which a plurality of binding agents are deposited onto a solid support surface in the form of an array or pattern, find use in a variety of applications, including drug screening, nucleicacid sequencing, mutation analysis, and the like. One important use of biochips is in the analysis of differential gene expression, where the expression of genes in different cells, normally a cell of interest and a control, is compared and any discrepancies in expression are identified. In such assays, the presence of discrepancies indicates a difference in the classes of genes expressed in the cells being compared.

In methods of differential gene expression, arrays find use by serving as a substrate to which is bound polynucleotide "probe" fragments. One then obtains "targets" from analogous cells, tissues or organs of a healthy and diseased organism. The targets are then hybridized to the immobilized set of polynucleotide "probe" fragments. Differences between the resultant hybridization patterns are then detected and related to differences in gene expression in the two sources.

A variety of different array technologies have been developed in order to meet the growing need of the biotechnology industry, as evidences by the extensive number of patents and references listed in the relevant literature section below.

Despite the wide variety of array technologies currently in preparation or available on the market, there is a continued need to identify new array devices to meet the needs of specific applications. Of particular interest
would be the development of an array capable of providing high throughput analysis of differential gene expression.

## Relevant Literature

Patents and patent applications describing arrays of biopolymeric compounds and methods for their fabrication include: $5,242,974 ; 5,384,261$; 5,405,783; 5,412,087; 5,424,186; 5,429,807; 5,436,327; 5,445,934; 5,472,672; 5,527,681; 5,529,756; 5,545,531; 5,554,501; 5,556,752; 5,561,071; 5,599,895; 5,624,711; 5,639,603; 5,658,734; WO 93/17126; WO 95/11995; WO 95/35505; EP 742 287; and EP 799897.

Patents and patent application describing methods of using arrays in various applications include: $5,143,854 ; 5,288,644 ; 5,324,633 ; 5,432,049$; 5,470,710; 5,492,806; 5,503,980; 5,510,270; 5,525,464; 5,547,839; 5,580,732; 5,661,028; WO 95/21265; WO 96/31622; WO 97/10365; WO 97/27317; EP 373 203; and EP 785280.

Other references of interest include: Atlas Human cDNA Expression Array I (April 1997) CLONTECHniques XII: 4-7; Lockhart et al., Nature Biotechnology (1996) 14: 1675-1680; Shena et al., Science (1995) 270: 467470; Schena et al., Proc. Nat'l Acad. Sci. USA (1996) 93: 10614-10619; Shalon et al., Genome Res. (1996) 6:639-645; Milosavlkevic et al., Genome Res. (1996) 6:132-141; Nguyen et al., Genomics (1995) 29:207-216; Piétu et al., Genome Res. (1996) 6:492-503; Zhao et al., Gene (1995) 166:207-213; Chalifour et al., Anal. Biochem. (1994) 216:299-304; Heller et al., Proc. Nat'l Acad. Sci. USA (1997) 94: 2150-2155; and Schena, M., BioAssays (1996) 18:427-431.

Any discussion of documents, acts, materials, devices, articles or the like which has been included in the present specification is solely for the purpose of providing a context for the present invention. It is not to be taken as an admission that any or all of these matters form part of the prior art base or were common general knowledge in the field relevant to the present invention as it existed in Australia before the priority date of each claim of this application.

## SUMMARY OF THE INVENTION

Arrays of polynucleotide spots stably associated with the surface of a solid support and kits comprising the same, as well as methods for their
preparation and use in hybridization assays, are provided. The subject arrays comprise a plurality of polynucleotide spots, wherein each different polynucleotide spot is made up of a polynucleotide probe composition and at least a portion of the polynucleotide probe compositions are made up of unique polynucleotides. The arrays are further characterized in that all of the unique polynucleotides on the array correspond to the same type of gene. The subject arrays find particular use in differential gene expression analysis. Also provided are sets of a representational number of gene specific primers useful in generating target nucleic acids for use with the subject arrays in hybridization assays.

Accordingly, the invention provides in a first aspect an array comprising a plurality of polynucleotide probe spots stably associated with the surface of a solid support, wherein each polynucleotide probe spot is made up of unique polynucleotides each having a length that does not exceed about 1000 nt, wherein each of said polynucleotides has a known sequence of a known gene whose coding region has been fully sequenced, where all of the unique polynucleotides on said array correspond to genes of a specific type.

In another aspect the invention provides a method of preparing an array according to the above said method comprising:
enzymatically generating said unique polynucleotides; and
stably associating said enzymatically-generated, complementary, unique polynucleotides on the surface of said solid support.

In another aspect the invention provides a composition that is a mixture of a representative number of distinct gene specific primers corresponding to at least twenty distinct genes.

In another aspect the invention provides a method for detecting expression of a gene using a hybridization assay, said method comprising:
contacting at least one labeled target polynucleotide sample with an array as previously described under hybridization conditions sufficient to produce a hybridization pattern; and
detecting said hybridization pattern.
In another aspect the invention provides a kit for use in a hybridization assay, said kit comprising:
an array as previously described.

In another aspect the invention provides a kit for use in detecting the differential expression of genes of a plurality of physiological sources, the kit comprising:
a set of a representative number of distinct gene specific primers as previously described.

Throughout this specification the word "comprise", or variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated element, integer or step, or group of elements, integers or steps, but not the exclusion of any other element, integer or step, or group of elements, integers or steps.

## BRIEF DESCRIPTION OF THE FIGURES

Fig. 1 provides a representation of an array according to the subject invention.

## DEFINITIONS

The term "nucleic acid" as used herein means a polymer composed of nucleotides, e.g. deoxyribonucleotides or ribonucleotides.

The terms "ribonucleic acid" and "RNA" as used herein mean a polymer composed of ribonucleotides.

The terms "deoxyribonucleic acid" and "DNA" as used herein mean a polymer composed of deoxyribonucleotides.

The term "oligonucleotide" as used herein denotes single stranded nucleotide multimers of from about 10 to 100 nucleotides in length.

The term "polynucleotide" as used herein refers to single or double stranded polymer composed of nucleotide monomers of greater than about 120 nucleotides in length up to about 1000 nucleotides in length.

The term "array type" refers to the type of gene represented on the array by the unique polynucleotides, where the type of gene that is represented on the array is dependent on the intended purpose of the array, e.g. to monitor expression of key human genes, to monitor expression of know oncogenes, etc, i.e. the use for which the array is designed. As such, all of the unique polynucleotides on a given array correspond to the same type or
category or group of genes. Genes are considered to be of the same type if they share some common linking characteristics, such as: species of origin, e.g. human, mouse, rat, etc.; tissue or cell type of origin, e.g. muscle. neural, dermal, organ, etc.; disease state, e.g. cancer; functions, e.g. protein kinases, tumor supressors and the like, participation in the same normal biological process, e.g. apoptosis, signal transduction, cell cycle regulation, proliferation, differentiation etc.; and the like. For example, one array type that is provided below is a "cancer array" in which each of the "unique" polynucleotide probes correspond to a gene associated with a cancer disease state. Likewise, a "human array" may be an array of polynucleotides corresponding to unique tightly regulated human genes. Similarly, an "apoptosis array" may be an array type in which the polynucleotides correspond to unique genes associated with apoptosis.

The "unique" polynucleotide sequences associated with each type of array of the present invention are sequences which are distinctive or different with respect to every other polynucleotide sequence on the array and correspond to the same type of gene, as defined above. For example, in a cancer array, each unique polynucleotide has a sequence that is not homologous to any other known cancer associated sequence. Moreover, each polynucleotide sequence on the array is statistically chosen to ensure that the probability of homology to any sequence of that type is very low. Morever, in the cancer array embodiment, all sequences are statistically chosen to insure that the probability of homology to any other sequence associated with cancer or of human origin is very low. An important feature of the individual polynucleotide probe compositions of the subject arrays is that they are only a fragment of the entire cDNA of the gene to which they correspond. In other words, for each gene represented on the array, the entire cDNA sequence the gene is not represented on the array. Instead, the sequence of only a portion or fragment of the entire cDNA is represented on the array by this unique polynucleotide.

The term "polynucleotide probe composition" refers to the nucleic acid composition that makes up each of the spots on the array. Thus, the term " polynucleotide probe composition" includes nucleic acid compositions of unique polynucleotides and control or calibrating polynucleotides (e.g. polynucleotides corresponding to housekeeping genes). The polynucleotide compositions are made up of single stranded polynucleotides (i.e. polynucleotides that are not hybridized to each other), where all of the polynucleotides in the probe composition may be identical to each other or there may be two different
polynucleotides (polynucleotides of different nucleotide sequence) in each probe composition, where the two different polynucleotides are complementary to each other.

The term "gene specific primer" means a polynucleotide of sufficient length to specifically hybridize to a distinct nucleic acid member of the sample, e.g. RNA or cDNA, where the length of the gene specific primers will usually be at least 8 nt . more usually at least 20 nt and may be as long as 25 nt or longer, but will usually not exceed 50 nt . The gene specific primers of the subject invention are sufficiently specific to hybridize to complementary template sequence during the generation of labeled nucleic acids under conditions sufficient for first strand cDNA synthesis, which conditions are known by those of skill in the art. The number of mismatches between the gene specific primer sequences and their complementary template sequences to which they hybridize during the generation of labeled nucleic acids in the subject methods will generally not exceed $20 \%$, usually will not exceed $10 \%$ and more usually will not exceed $5 \%$, as determined using the FASTA program using default settings.

## DESCRIPTION OF THE SPECIEIC EMBODIMENTS

Arrays of polynucleotide spots and methods for their preparation are provided. In the subject arrays, a plurality of polynucleotide spots is stably associated with the surface of a solid support, where at least a portion of the polynucleotide spots on the array are made up of unique polynucleotides and all of the unique polynucleotides of the array correspond to one particular type of gene, e.g. tightly regulated human genes, genes associated with a particular disease state, genes associated with cell cycle regulation, etc. The subject arrays find particular use in gene expression assays. Also provided are sets of a representational number of gene specific primers useful in generating target nucleic acids for use with the subject arrays. In further describing the subject invention. the arrays first will be described in general terms. Next, methods for their preparation are described. Following this, a description of representative specific array types falling within the scope of the invention will be provided. Finally, a review of representative applications in which the subject arrays may be employed will be provided, where this review includes a description of the sets of a representational number of gene specific primers according to the subject invention.

Before the subject invention is further described, it is to be understood that the invention is not limited to the particular embodiments of the invention described below, as variations of the particular embodiments may be made and still fall within the scope of the appended claims. It is also to be understood that the terminology employed is for the purpose of describing particular embodiments, and is not intended to be limiting. Instead, the scope of the present invention will be established by the appended claims.

In this specification and the appended claims, the singular forms "a," "an," and "the" include plural reference unless the context clearly dictates otherwise. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood to one of ordinary skill in the art to which this invention belongs.

## Arrays of the Subject Invention-General Description

## Array Structure

The arrays of the subject invention have a plurality of polynucleotide spots stably associated with a surface of a solid support. Each spot on the array comprises a polynucleotide sample, i.e. polynucleotide probe composition, of known identity, usually of known sequence, as described in greater detail below. The polynucleotide spots on the array may be any convenient shape, but will typically be circular, elliptoid, oval or some other analogously curved shape. The density of the spots on the solid surface is at least about $5 / \mathrm{cm}^{2}$ and usually at least about $10 / \mathrm{cm}^{2}$ but does not exceed about $1000 / \mathrm{cm}^{2}$, and usually does not exceed about $500 / \mathrm{cm}^{2}$, and more usually does not exceed about $300 / \mathrm{cm}^{2}$. The spots may be arranged in any convenient pattern across or over the surface of the array, such as in rows and columns so as to form a grid, in a circular pattern, and the like, where generally the pattern of spots will be present in the form of a grid across the surface of the solid support. See Fig. 1.

In the subject arrays, the spots of the pattern are stably associated with the surface of a solid support, where the support may be a flexible or rigid solid support. By stably associated is meant that the polynucleotides of the spots maintain their position relative to the solid support under hybridization and washing conditions. As such, the polynucleotide members which make up the spots can be non-covalently or covalently stably associated
with the support surface. Examples of non-covalent association include non-specific adsorption. binding based on electrostatic (e.g. ion. ion pair interactions), hydrophobic interactions, hydrogen bonding interactions, specific binding through a specific binding pair member covalently attached to the support surface. and the like. Examples of covalent binding include covalent bonds formed between the spot polynucleotides and a functional group present on the surface of the rigid support, e.g. - OH , where the functional group may be naturally occurring or present as a member of an introduced linking group, as described in greater detail below.

As mentioned above, the array is present on either a flexible or rigid substrate. By flexible is meant that the support is capable of being bent, folded or similarly manipulated without breakage. Examples of solid materials which are flexible solid supports with respect to the present invention include membranes, e.g. nylon, flexible plastic films, and the like. By rigid is meant that the support is solid and does not readily bend, i.e. the support is not flexible. As such, the rigid substrates of the subject arrays are sufficient to provide physical support and structure to the polymeric targets present thereon under the assay conditions in which the array is employed, particularly under high throughput handling conditions. Furthermore, when the rigid supports of the subject invention are bent, they are prone to breakage.

The solid supports upon which the subject patterns of spots are presented in the subject arrays may take a variety of configurations ranging from simple to complex, depending on the intended use of the array. Thus, the substrate could have an overall slide or plate configuration, such as a rectangular or disc configuration. In many embodiments, the substrate will have a rectangular cross-sectional shape, having a length of from about 10 mm to 200 mm , usually from about 40 to 150 mm and more usually from about 75 to 125 mm and a width of from about 10 mm to 200 mm , usually from about 20 mm to 120 mm and more usually from about 25 to 80 mm , and a thickness of from about 0.01 mm to .5 .0 mm , usually from about 0.1 mm to 2 mm and more usually from about 0.2 to 1 mm .

The substrates of the subject arrays may be fabricated from a variety of materials. The materials from which the substrate is fabricated should ideally exhibit a low level of non-specific binding during hybridization events. In many situations, it will also be preferable to employ a material that is transparent to visible and/or UV light. For flexible substrates, materials of interest include: nylon, both modified and unmodified, nitrocellulose.
polypropylene. and the like, where a nylon membrane, as well as derivatives thereof, is of particular interest in this embodiment. For rigid substrates, specific materials of interest include: glass; plastics, e.g. polytetrafluoroethylene, polypropylene, polystyrene, polycarbonate, and blends thereof, and the like; metals, e.g. gold, platinum, and the like; etc.

The substrates of the subject arrays comprise at least one surface on which the pattern of spots is present, where the surface may be smooth or substantially planar, or have irregularities, such as depressions or elevations. The surface on which the pattern of spots is present may be modified with one or more different layers of compounds that serve to modify the properties of the surface in a desirable manner. Such modification layers, when present, will generally range in thickness from a monomolecular thickness to about 1 mm , usually from a monomolecular thickness to about 0.1 mm and more usually from a monomolecular thickness to about 0.001 mm . Modification layers of interest include: inorganic and organic layers such as metals, metal oxides, polymers, small organic molecules and the like. Polymeric layers of interest include layers of: peptides, proteins, polynucleic acids or mimetics thereof, e.g. peptide nucleic acids and the like; polysaccharides, phospholipids, polyurethanes, polyesters, polycarbonates, polyureas, polyamides, polyethyleneamines, polyarylene sulfides, polysiloxanes, polyimides, polyacetates, and the like, where the polymers may be hetero- or homopolymeric, and may or may not have separate functional moieties attached thereto, e.g. conjugated.

The total number of spots on the substrate will vary depending on the number of different polynucleotide probes one wishes to display on the surface, as well as the number of control spots, calibrating spots and the like, as may be desired depending on the particular application in which the subject arrays are to be employed. Generally, the pattern present on the surface of the array will comprise at least about 10 distinct spots, usually at least about 20 distinct spots, and more usually at least about 50 distinct spots, where the number of spots may be as high as 10,000 or higher, but will usually not exceed about 5.000 distinct spots, and more usually will not exceed about 3,000 distinct spots. In many embodiments, it is preferable to have each distinct probe composition presented in duplicate, i.e. so that there are two spots for each distinct polynucleotide probe composition of the array. In certain embodiments, the number of spots will range from about 200 to 600 .

The amount of polynucleotide present in each spot will be sufficient to provide for adequate hybridization and detection of target nucleic acid during the assay in which the
array is employed. Generally, the amount of polynucleotide in each spot will be at least about 0.1 ng , usually at least about 0.5 ng and more usually at least about 1 ng , where the amount may be as high as 1000 ng or higher, but will usually not exceed about 20 ng and more usually will not exceed about 10 ng . The copy number of each polynucleotide in a spot will be sufficient to provide enough hybridization sites for target molecule to yield a detectable signal, and will generally range from about 0.01 fmol to 50 fmol , usually from about 0.05 fmol to 20 fmol and more usually from about 0.1 fmol to 5 fmol . Where the spot has an overall circular dimension, the diameter of the spot will generally range from about 10 to $5,000 \mu \mathrm{~m}$, usually from about 20 to $2,000 \mu \mathrm{~m}$ and more usually from about 50 to 1000 $\mu \mathrm{m}$.

A critical feature of the subject arrays is that at least a portion, usually the majority, of the polynucleotide spots on the array are made up of polynucleotide probes that all correspond to the same kind or kind of gene, i.e. genes that all share some common characteristic or can be grouped together based on some common feature, such as species of origin, tissue or cell of origin, functional role, disease association, etc. Other spots which may be present in the pattern include spots comprising genomic DNA, housekeeping genes, negative and positive control genes, and the like. These latter types of spots comprise polynucleotides that are not "unique" as that term is defined and used herein, i.e. they are "common." In other words, they are calibrating or control genes whose function is not to tell whether a particular "key" gene of interest is expressed, but rather to provide other useful information, such as background or basal level of expression, and the like. The percentage of spots which are made of unique polynucleotides that correspond to the same type of gene is generally at least about 30 number $\%$, and usually at least about 60 number $\%$ and more usually at least about 80 number $\%$. Therefore, the arrays of the subject invention will be of a specific array type, where representative array types include: human arrays, mouse arrays, cancer arrays, apoptosis arrays, human stress arrays, oncogene and tumor suppressor arrays, cell-cell interaction arrays, cytokine and cytokine receptor arrays, rat arrays, blood arrays, mouse stress arrays, neuroarrays, and the like, where some of these representative arrays are described in greater detail below.

With respect to the polynucleotide probes that correspond to a particular type or kind of gene, type or kind can refer to a plurality of different characterizing features, where such features include: species specific genes, where specific species of interest include eukaryotic
species, such as mice, rats, rabbits, pigs, primates, humans, etc.; function specific genes, where such genes include oncogenes, apoptosis genes, cytokines, receptors. protein kinases, etc.; genes specific for or involved in a particular biological process, such as apoptosis, differentiation, cell cycle regulation, cancer, aging, proliferation, etc.; location specific genes. where locations include organ, such as heart, liver, prostate, lung etc., tissue, such as nerve, muscle, connective, etc., cellular, such as axonal, lymphocytic, etc, or subcellular locations, e.g. nucleus, endoplasmic reticulum, Golgi complex, endosome, lyosome, peroxisome, mitochondria, cytoplasm, cytoskeleton, plasma membrane, extracellular space; specific genes that change expression level over time, e.g. genes that are expressed at different levels during the progression of a disease condition, such as prostate genes which are induced or repressed during the progression of prostate cancer.

The average length of the polynucleotides on the array is chosen to be of sufficient length to provide a strong and reproducible signal, as well as tight and robust hybridization. As such, the average length of the polynucleotides of the array will typically range from about 120 to 1000 nt and usually from about 120 to 800 nt , where in many embodiments, the average length ranges from about 200 to 700 nt , and usually 200 to 600 nt . The length of each polynucleotide on the array is less than the length of the mRNA to which it corresponds. As such, the polynucleotide represents only a fraction of the full length cDNA to which it corresponds.

As mentioned above, the subject arrays typically comprise one or more additional spots of polynucleotides which do not correspond to the array type, i.e. the type or kind of gene represented on the array. In other words, the array may comprise one or more spots that are made of non "unique" polynucleotides, i.e common polynucleotides. For example, spots comprising genomic DNA may be provided in the array, where such spots may serve as orientation marks. Spots comprising plasmid and bacteriophage genes, genes from the same or another species which are not expressed and do not cross hybridize with the cDNA target, and the like, may be present and serve as negative controls. In addition, spots comprising housekeeping genes and other control genes from the same or another species may be present, which spots serve in the normalization of mRNA abundance and standardization of hybridization signal intensity in the sample assayed with the array.

## Polynucleotide Probes of the Arrays:

Each spot of the pattern present on the surface of the substrate is made up of a unique polynucleotide probe composition. By "polynucleotide probe composition" is meant a collection or population of single stranded polynucleotides capable of participating in a hybridization event under appropriate hybridization conditions, where each of the individual polynucleotides may be the same -- have the same nucleotide sequence-- or different sequences, for example the probe composition may consist of 2 different single stranded polynucleotides that are complementary to each other (i.e. the two different polynucleotides in the spot are complementary but physically separated so as to be single stranded, i.e. not hybridized to each other). In many embodiments, the probe compositions will comprise two complementary, single stranded polynucleotides.

In those polynucleotide probe compositions having unique polynucleotides, the sequence of the polynucleotides are chosen in view of the type and the intended use of the array on which they are present. The unique polynucleotides are chosen so that each distinct unique polynucleotide does not cross-hybridize with any other distinct unique polynucleotide on the array, i.e. the polynucleotide of any other polynucleotide probe composition that corresponds to a different gene falling within the broad category or type of genes represented on the array. As such, the nucleotide sequence of each unique polynucleotide of a probe composition will have less than $90 \%$ homology, usually less than $85 \%$ homology, and more usually less than $80 \%$ homology with any other different polynucleotide of a probe composition of the array, where homology is determined by sequence analysis comparison using the FASTA program using default settings. The sequence of unique polynucleotides in the probe compositions are not conserved sequences found in a number of different genes (at least two), where a conserved sequence is defined as a stretch of from about 40 to 200 nucleotides which have at least about $90 \%$ sequence identity, where sequence identity is measured as above. The polynucleotide will generally be a deoxyribonucleic acid having a length of from about 120 to 1000 , usually from 120 to 700 nt . and more usually 200 to 600 $n t$. The polynucleotide will not cross-hybridize with any other polynucleotide on the array under standard hybridization conditions. Again, the length of the polynucleotide will be shorter than the mRNA to which it corresponds.

## Array Preparation

The subject arrays can be prepared using any convenient means. One means of preparing the subject arrays is to first synthesize the polynucleotides for each spot and then deposit the polynucleotides as a spot on the support surface. The polynucleotides may be prepared using any convenient methodology, such as automated solid phase synthesis protocols. preparative PCR and like, where preparative PCR or enzymatic synthesis is preferred in view of the length and the large number of polynucleotides that must be generated for each array.

For preparative $\operatorname{PCR}$, primers flanking either side of the portion of the gene of interest will be employed to produce amplified copy numbers of the portion of interest. Methods of performing preparative PCR are well known in the art, as summarized in PCR, Essential Techniques (Ed. J.F. Burke, John Wiley \& Sons)(1996). Alternatively, if a gene fragment of interest is cloned into a vector, vector primers can be used to amplify the gene fragment of interest to produce the polynucleotide.

In determining the portion of the gene to be amplified and subsequently placed on the array, regions of the gene having a sequence unique to that gene should preferably be amplified. Different methods may be employed to choose the specific region of the gene to be amplified. Thus, one can use a random approach based on availability of a gene of interest. However, instead of using a random approach which is based on availability of a gene of interest, a rational design approach may also be employed to choose the optimal sequence for the hybridization array. Preferably, the region of the gene that is selected and amplified is chosen based on the following criteria. First, the sequence that is chosen should yield a polynucleotide that does not cross-hybridize with any other polynucleotide that is present on the array. Second, the sequence should be chosen such that the polynucleotide has a low probability of cross-hybridizing with a polynucleotide having a nucleotide sequence found in any other gene, whether or not the gene is to be represented on the array from the same species of origin, e.g. for a human array, the sequence will not be homologous to any other human genes. As such, sequences that are avoided include those found in: highly expressed gene products, structural RNAs, repeated sequences found in the sample to be tested with the array and sequences found in vectors. A further consideration is to select sequences which provide for minimal or no secondary structure, structure which allows for
optimal hybridization but low non-specific binding, equal or similar thermal stabilities, and optimal hybridization characteristics.

The prepared polynucleotides may be spotted on the support using any convenient methodology, including manual techniques, e.g. by micro pipette, ink jet, pins, etc., and automated protocols. Of particular interest is the use of an automated spotting device, such as the Beckman Biomek 2000 (Beckman Instruments). As mentioned above, the polynucleotide probe compositions that are spotted onto the array surface are made up of single stranded polynucleotides, where all the polynucleotides may be identical to each other or a population of complementary polynucleotides may be present in each spot.

## Specific Array Types of the Subject Invention

A variety of specific array types are also provided by the subject invention. As discussed above, array type refers to the nature of the polynucleotide probes present on the array and the types of genes to which the probes correspond. These array types include: human array; mouse array; cancer array, apoptosis array, human stress array, oncogene and tumor suppressor arrray, cell-cell interaction array, and cytokine and cytokine receptor array, as well as other types of arrays, e.g. rat array, rat stress array, blood array, mouse stress array, and nueroarray. Each of these arrays is described separately below.

## Human Array

One specific array type provided by the subject invention is the human array. In the human array of the subject invention, the majority of the spots on the array have a polynucleotide sequence corresponding to a human gene of interest. As such, all of the unique polynucleotide probes on the array correspond to human genes. The human genes represented on the human array are typically those genes that have been identified by those of skill in the art as key genes. By "key" is meant that the genes are relevant and related to the purpose of the array, e.g. the identification of difference in the expression profiles of different cell or tissue types, where the key genes are generally functionally important to the cell. In many embodiments, the genes represented on the human array are tightly regulated human genes. The term "tightly regulated gene" is used herein in accordance with its art accepted definition and use. As such, by tightly regulated human gene is meant a gene which
is not "leaky," as opposed to housekeeping genes which are generally expressed at similar levels in different cells and different tissues, i.e. a gene which is inducible such that in response to a specific inducing signal the gene turns "on" and when this signal is removed, the gene turns "off."

In certain embodiments of the human array, human genes that may be represented on the subject arrays include: (a) oncogenes \& tumor suppressors; (b) cell cycle regulators; (c) stress response proteins; (d) ion channel \& transport proteins; (e) intracellular signal transduction modulators and effectors; (f) apoptosis-related proteins; (g) DNA synthesis, repair and recombination proteins; (h) transcription factors \& general DNA binding proteins; (i) growth factor \& chemokine receptors; (j) interleukin \& interferon receptors; (k) hormone receptors; (l) neurotransmitter receptors; (m) cell surface antigens \& cell adhesion proteins; (n) growth factors, cytokines and chemokines; (o) interleukins \& interferons; (p) hormones; (q) extracellular matrix proteins; (r) cytoskeleton \& motility proteins; (s) RNA processing \& turnover proteins; (t) post-translational modification, trafficking \& targeting proteins; (u) protein turnover; and (v) metabolic pathway proteins.

In view of the length of the polynucleotides of the probe compositions of the spots, each polynucleotide of a probe composition typically has a nucleotide sequence of only a portion of the human gene. Specific sequences to which the polynucleotide sequence may correspond include those identified in Table 1 below, where by "correspond" is meant that the polynucleotide could have the same sequence as specified or a sequence complementary to the specified sequence. Whether the polynucleotide sequence is the same as a portion of the sense strand of the gene to which is corresponds or complementary thereto is based primarily on the nature of the target which the array is to be used, e.g. if the target is first strand cDNA, the polynucleotide will have a sequence found in the anti-sense DNA strand of the gene to which it corresponds.

Of particular interest is a human array of the subject invention as shown in Fig. 1. In the array, each spot on the array comprises a known polynucleotide, as specified in Table 1, where the array comprises spots which: (a) correspond to 588 different tightly regulated human genes; (b) comprise plasmid and bacteriophage polynucleotides; (c) comprise polynucleotides corresponding to housekeeping genes: and (d) genomic DNA. Each of the different types of polynucleotide spots are positioned at a known location on the membrane surface.


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| F1h | $\times 04571$ | kidney epidermal growth factor (EGF) precursor | 4164-4434 |  |
| E3a | J03171 | interferon alpha receptor (HUIFN-ALPHA-REC) | 2562-2740 |  |
| F6i | M57627 | interleukin 10 (IL10) | 442-648 |  |
| E3b | M26062 | interleukin 2 receptor beta chain (P70-75) | 3399-3748 |  |
| E3c | M74782 | interleukin 3 receptor (HIL-3RA) | 651-1116 |  |
| E3d | X52425 | interleukin 4 receptor | 2641-2974 |  |
| E3e | M75914 | interleukin 5 receptor alpha | 555-959 |  |
| E3f | X77722 | interferon alpha/beta receptor | 553-1012 |  |
| Fli | HG1621 | cytokine humig | 2021-2246 |  |
| E4g | HG1160, M37981 | cholinergic receptor nicotinic alfa polipeptide 3 | 934-1250 |  |
| E3g | HG1252, D11086 | interleukin 2 receptor gamma polipeptide | 674-1006 |  |
| E4b | HG1334, M20132, J03180 | androgen receptor | 1879-2146 |  |
| E1b | HG135, M73238 | ciliary neurotropic factor receptor | 610-849 |  |
| C1h | HG1410, X68486 | adenosine receptor | 1281-1494 |  |
| E3h | HG1757, J03143 | interferon gamma receptor | 610-824 |  |
| E1c | HG2246, M60459 | erythropoietin receptor | 1423-1740 |  |
| $\overline{\mathrm{C} 1 \mathrm{i}}$ | 556143 | A1 adenosine receptor-adenylate cyclase inhibitor | 508-921 |  |
| B1e | HG3354, Z30425 | orphan hormone nuclear receptor | 817-1147 |  |
| C1j | HG3381, X76981 | adenosine receptor A3 | 1043-1452 |  |
| E 4 c | L00587 | calcitonin receptor | 885-1270 |  |
| B1f | HG74, M62424 | coagulating factor II receptor | 2297-2697 |  |
| A1e | HG886, L07594 | transforming growth factor beta receptor III 300 kDa | 3358-3592 |  |
| Ē3i | HG216, M84747 | interleukin 9 receptor | 289-528 |  |
| E3j | HG4080, U00672 | interleukin 10 receptor | 2448-2803 |  |
| E1d | HG423, M14764 | nerve growth factor receptor | 2762-3242 |  |
| E5d | HG1023 | Vitronectin receptor alpha subunit | 2442-2473 |  |
| D1b | HG125 | GATA-binding protein 2 | 1126-1363 |  |
| D1c | HG1377 | CCAAT-box DNA-binding protein Hap2 homolog | 958-1272 |  |
| C1k | HG1458 | retinoic acid receptor epsilon | 1315-1633 |  |
| A1f | HG1470, X13293 | B-myb | 1873-2272 |  |
| B1g | HG1551 | tyrosine kinase receptor tie | 3114-3536 |  |
| C11 | HG1601 | tyrosine kinase receptor FLT4 class III | 4236-4402 |  |
| D1d | HG1603 | helix-loop-helix protein 1R21 | 858-560 |  |
| F1j | HG1650 | thrombomodulin | 1262-1605 |  |
| D1e | HG1697 | basic transcription element-binding protein 2 | 572-976 |  |
| D11 | HG1963 | basic transcription factor 62 kDa subunit | 1449-1831 |  |



| Array Coordinate | GeneBank\# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| B2c | HG918 | tyrosine phosphatase receptor gamma polypeptide | 3623-3938 |  |
| D2h | HG970 | DNA-binding protein PO-GA | 3196-3413 |  |
| D2i | HG99, M64673 | CCAAT enhancer-binding protein beta | 294-572 |  |
| A11 | J04111 | c-jun proto-oncogene (jun) clone $\mathrm{HCJ}-1$ | 2207-2583 |  |
| E31 | M27492 | interleukin 1 receptor | 3847-4288 |  |
| C1m | M 33294 | tumor necrosis factor receptor | 1570-1817 |  |
| F1m | M37435 | macrophage-specific colony-stimulating factor (CSF-1) | 2277-2413 |  |
| A1m | YOO285 | insulin-like growth factor II receptor | 1394-1831 |  |
| $\overline{\text { A1n }}$ | HG404 | tyrosine kinase receptor HER2 | 2556-2722 |  |
| B2d | D10923 | HM74 | 1357-1826 |  |
| B2e | D10924 | HM89 | 351-808 |  |
| B21 | D10925 | HM145 | 1353-1832 |  |
| F1n | D14012 | hepatocyte growth factor activator precursor | 1487-1845 |  |
| F2a | D16431 | hepatoma-derived growth factor | 359-625 |  |
| F2b | D30751 | bone morphogenetic protein 4 | 943-1321 |  |
| B2g | J03358 | FER tyrosine kinase | 2384-2688 |  |
| F2c | J04130 | activation (Act-2) | 236-592 |  |
| F2d | J05081 | endothelin ET3 | 1428-1685 |  |
| F2e | K03515 | neuroleukin | 1368-1656 |  |
| Ā2a | L06139 | TEK tyrosine kinase receptor | 3243-3586 |  |
| E19 | L06622 | endothelin receptor EDNRA | 870-1080 |  |
| E1h | L06623 | endothelin receptor EDNRB | 497-814 |  |
| F6g | L06801 | interleukin IL-13 | 285-743 |  |
| C1n | L07414 | CD40 ligand | 863-1277 |  |
| C2a | L08096 | CD27 ligand | 233-627 |  |
| E3m | L08187 | cytokine receptor (EB13) | 627-1019 |  |
| F2f | L12260 | glial growth factor 2 (recombinant) | 1069-1452 |  |
| F2g | L12261 | glial growth factor (recombinant) | 762-1041 |  |
| F6h | L15344 | interleukin IL-14 | 1181-1562 |  |
| F2h | L36052 | thrombopoietin (MGDF/Mpl ligand) | 230-613 |  |
| E1i | M10051 | insulin receptor | 3274-3758 |  |
| F2i | M17778 | uromodulin | 1463-1913 |  |
| F2j | M21121 | RANTES pro-inflammatory cytokine | 180-545 |  |
| E1j | M21574 | PDGF-alpha receptor | 5118-5583 |  |
| E1k | M21616 | PDGF-beta receptor | 842-1133 |  |
| F2k | M22488 | bone morphogenetic protein 1 | 702-1098 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| F21 | M22489 | bone morphogenetic protein 2a | 567-997 |  |
| F2m | M22491 | bone morphogenetic protein 3 | 1458-1731 |  |
| F2n | M23452 | macrophage inflammatory protein GOS19-1 | 243-704 |  |
| F3a | M24545 | monocyte chemotactic and activating factor MCAF | 36-384 |  |
| F3b | M25667 | neuronal growth protein GAP-43 | 747-1154 |  |
| F3c | M27288 | oncostatin M | 833-1113 |  |
| F3d | M30704 | amphiregulin AR | 511.837 |  |
| F3e | M31145 | insuline-like growth factor binding protein 1 | 476-861 |  |
| E11 | M31165 | TNF-inducible hyaluronate-binding protein TSG-6 | 320-584 |  |
| F31 | M32977 | heparin-binding vascular endothelial growth factor VEGF | 198-622 |  |
| A2b | M35410 | insuline-like growth factor binding protein 2 | 680-1071 |  |
| F7a | M36717 | ribonuclease/angiogenun inhibitor RAI | 713-1028 |  |
| F3g | M37722 | bFGF receptor | 1746-1967 |  |
| B2h | M57230 | glycoprotein gp130 | 1757-2152 |  |
| F3h | M57399 | nerve growth factor HBNF-1 | 602-847 |  |
| F3i | M57502 | secreted protein 1-309 | 205-397 |  |
| F $\overline{6} \mathrm{i}$ | M57765 | interleukin IL-11 | 132-460 |  |
| E1m | M59818 | granulocyte colony-stimulating factor receptor G-CSFR1 | 1453-1891 |  |
| F3j | M59964 | stem cell factor | 898-1283 |  |
| F3k | M60278 | heparin-binding EGF-like growth factor | 1905-2146 |  |
| F31 | M60718 | HGF (hepatocyte growth factor) | 1549-1970 |  |
| F3m | M60828 | keratinocyte growth factor | 419-766 |  |
| F3n | M61176 | brain-derived neurotrophic factor BDNF | 982-1265 |  |
| F4a | M62302 | growth/differentiation factor GDF-1 | 615-957 |  |
| E1n | M62505 | C5a anaphylatoxin receptor | 725-1098 |  |
| E5e | M63928 | T cell activation antigen CD27 | 513-977 |  |
| F4b | M65199 | endothelin ET2 | 338-570 |  |
| F6j | M65290 | interleukin IL-12 (NKSF p40) | 622-848 |  |
| F6k | M65291 | interleukin IL-12 (NKSF p35) | 600-990 |  |
| C2b | M67454 | Fas antigen | 2063-2288 |  |
| E2a | M68932 | interleukin 8 receptor alpha (IL8RA) | 1179-1370 |  |
| E2b | M73482 | NMB-R (neuromedin B receptor) | 282-544 |  |
| F4C | M74178 | hepatocyte growth factor-like protein | 1643-2015 |  |
| $\overline{\text { A5 }}$ | M76125 | AXL tyrosine kinase receptor | 2054-2328 |  |
| E5 5 | M83554 | lymphocyte activation antigen CD30 | 3152-3421 |  |
| F4̈d | M92381 | thymosin beta-10 | 40-342 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| F4e | M92934 | connective tissue growth factor | 1459-1748 |  |
| C2c | M93426 | tyrosine phosphatase receptor zeta-polypeptide | 5090-1748 |  |
| F4i | M96956 | TDGF3 | 1294-1712 |  |
| E2c | S59184 | RYK=related to receptor tyrosine kinase isolog | 1760-1968 |  |
| A2c | U01134 | VEGF receptor | 1288-1604 |  |
| E2d | U01839 | Duffy blood group antigen (Fya-b+) | 127-150 |  |
| A5d | U02687 | growth factor receptor tyrosine kinase STK-1 | 2491-2965 |  |
| E3n | U03187 | interleukin 12 receptor component | 1053-1381 |  |
| E2e | U03882 | monocyte chemoattractant protein 1 receptor (MCP-1RA) alternatively spliced | 1514-1799 |  |
| E2f | U03905 | monocyte chemoattractant protein 1 receptor (MCP-1RB) alternatively spliced | 1362-1713 |  |
| C2d | U04806 | FLT3/FLK2 ligand | 29-362 |  |
| F4g | U10117 | endothelial-monocyte activating polypeptide II | 272-304 |  |
| E2g | U11814 | keratinocyte growth factor receptor | 753-1189 |  |
| C2e | U13737 | cysteine protease CPP32 isom alpha | 2007-2434 |  |
| F61 | U14407 | interleukin IL-15 | 338-695 |  |
| E2h | U14722 | activin type I receptor | 333.740 |  |
| F4h | U43142 | VRP (vascular endothelial growth factor related protein) | 1165-1559 |  |
| F4i | X02530 | IFN-gamma-inducible chemokine IP-10 | 280-613 |  |
| A1d | X06182 | c-kit proto-oncogene | 37-430 |  |
| F4i | $\times 06233$ | MRP-14 (calcium binding protein in macrophages MIF-related) | 16-254 |  |
| F4k | X06234 | MRP-8 (calcium binding protein in macrophages MIF-related) | 37-351 |  |
| F41 | X06374 | platelet-derived growth factor A chain PDGF-A | 522-955 |  |
| F4m | X13967 | leukemia inhibitory factor LIF | 1810-2239 |  |
| F6m | X17543 | interleukin IL-9 (P40) | 156-186 |  |
| E2i | X17648 | granulocyte-macrophage colony-stimulating factor receptor GM-CSFRa | 868-1173 |  |
| F4n | X51943 | fibroblast growth factor FGF-1 | 1131-1502 |  |
| F5a | X53655 | nerve growth factor NGF-2 (same as NT-3) | 112-416 |  |
| F5b | X53799 | macrophage inflammatory protein-2alpha (MIP2alpha) | 157-501 |  |
| F5c | X54936 | PIGF (placenta growth factor) | 1098-1371 |  |
| E4a | X59770 | interleukin 1 receptor type II | 842-1244 |  |
| E2j | X60592 | Cdw40 | 198-605 |  |
| Ē2k | X72304 | beta-thromboglobulin-like protein | 230-533 |  |
| F5d | X 78686 | neutrophil-activating peptide ENA-78 | 65-329 |  |
| F5e | X79929 | OX40 ligand/gp34 | 329-657 |  |



| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| Ḋ3g | L19606 | paired box homeotic protein (PAX8) | 113-338 |  |
| C5m | L20046 | ERCC5 excision repair protein | 1374-1638 |  |
| B3b | L20320 | protein serine/threonine kinase stk1 | 89-305 |  |
| B3C | L20321 | protein serine/threonine kinase stk2 | 2534-2802 |  |
| B3d | L20422 | 14-3-3n protein | 163-671 |  |
| D3h | L20433 | octamer binding transcription factor 1 (OTF1) | 3275-3583 |  |
| E5j | L20815 | S protein | 1677-2107 |  |
| B1a | L20977 | plasma membrane calcium ATPase isoform 2 (ATP2B2) | 3861-4236 |  |
| B3e | L22075 | guanine nucleotide regulatory protein (G13) | 1073-1376 |  |
| C2h | L22474 | Bax beta | 227-278 |  |
| C5n | L24564 | Rad | 489.780 |  |
| B3i | L24959 | calcium/calmodulin dependent protein kinase | 969-1220 |  |
| B3g | L25259 | CTLA4 counter-receptor (B7-2) | 496-722 |  |
| C2i | L29511 | GRB2 isoform | 355-573 |  |
| D3i | L31881 | nuclear factor 1-X | 415-729 |  |
| B3h | L32976 | protein kinase (MLK-3) | 970-1283 |  |
| A5g | L33264 | CDC2-related protein kinase (PISSLRE) | 454.755 |  |
| D3j | L34587 | RNA polymerase II elongation factor SIII p15 subunit | 115-354 |  |
| B3i | L35233 | autocrine motility factor receptor (AMFR) | 1221-1514 |  |
| $\overline{\text { A2h }}$ | M13150 | mas proto-oncogene | 262-726 |  |
| D3k | M14631 | guanine nucleotide-binding protein G-s alpha subunit partial cds | 824-1120 |  |
| B1b | M15800 | MAL protein | 461-695 |  |
| D31 | M16937 | homeobox c1 protein | 367-667 |  |
| E5k | M21097 | differentiation antigen (CD19) | 740-1071 |  |
| B3j | M22199 | protein kinase C alpha-polypeptide (PKCA) | 767-1106 |  |
| E51 | M23197 | differentiation antigen (CD33) | 885-1141 |  |
| A5h | M26708 | prothymosin alpha (ProT-alpha) | 538-864 |  |
| B3k | M28210 | GTP-binding protein (RAB3A) | 288-591 |  |
| B31 | M28211 | GTP-binding protein (RAB4) | 255-495 |  |
| B3m | M28212 | GTP-binding protein (RAB6) | 59-310 |  |
| B3n | M28213 | GTP-binding protein (RAB2) | 56-269 |  |
| B4a | M28214 | GTP-binding protein (RAB3B) | 322-621 |  |
| B4b | M28215 | GTP-binding protein (RAB5) | 447-672 |  |
| A5i | M28882 | MUC18 glycoprotein | 1756-2180 |  |
| $\overline{\mathrm{D}} 3 \mathrm{~m}$ | M29038 | stem cell protein (SCL) | 2804-3086 |  |
| A $\overline{5}$ j | M29142 | myeloblastin | 312-693 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| Ė5m | M30257 | vascular cell adhesion molecule 1 | 1056-1450 |  |
| E5n | M30640 | endothelial leucocyte adhesion molecule I (ELAM1) | 2098-2549 |  |
| C6a | M30938 | Ku (p70/p80) subunit | 2340-2764 |  |
| Ā2i | M31213 | papillary thyroid carcinoma-encoded protein | 2285-2631 |  |
| D3n | M31523 | transcription factor (E2A) | 2277-2685 |  |
| B4c | M31630 | cyclic AMP response element-binding protein (HB16) 3' end | 316-636 |  |
| C6b | M31899 | DNA repair helicase (ERCC3) | 2109-2466 |  |
| C6c | M32865 | Ku protein subunit | 1729-1974 |  |
| E6a | M33374 | cell adhesion protein (SQM1) | 53-354 |  |
| E6b | M34064 | N -cadherin | 942-1299 |  |
| B4d | M34356 | active transcription factor CREB | 433-780 |  |
| D4a | M34960 | transcription factor IID | 561-843 |  |
| C6d | M36089 | DNA-repair protein (XRCC1) | 1226-1539 |  |
| B4e | M36429 | transducin beta-2 subunit | 443-789 |  |
| B41 | M36430 | transducin beta-1 subunit 3' end | 58-338 |  |
| D4b | M36542 | lymphoid-specific transcription factor | 647-942 |  |
| D4c | M36711 | sequence-specific DNA-binding protein (AP-2) | 950-1211 |  |
| A2j | M54915 | h -pim-1 protein (h-pim-1) | 893-1187 |  |
| E6c | M54992 | B cell differentiation antigen | 963-1224 |  |
| E6d | M59040 | cell adhesion molecule (CD44) | 1158-1408 |  |
| A2k | M60915 | neurofibromatosis protein type I (NF1) | 740-1027 |  |
| D4d | M62397 | colorectal mutant cancer protein | 3626-3902 |  |
| D4e | M62810 | mitochondrial transcription factor 1 | 640-668 |  |
| D4f | M62829 | transcription factor ETR103 | 989-1276 |  |
| D4g | M62831 | transcription factor ETR101 | 1018-1410 |  |
| C6e | M63488 | replication protein A 70kDa subunit | 1498-1838 |  |
| A5k | M63618 | bullous pemphigoid antigen | 5680-6055 |  |
| D4h | M63896 | transcriptional enhancer factor (TEF 1) DNA | 2935-3238 |  |
| E6e | M74387 | cell adhesion molecule L1 (L1CAM) | 3197-3483 |  |
| C6i | M74524 | HHR6A (yeast RAD 6 homologue) | 175-433 |  |
| E6f | M74777 | dipeptidyl peptidase IV (CD26) | 1205-1507 |  |
| C̄2j | M74816 | sulfated glycoprotein-2 3'end | 709-990 |  |
| D4i | M75952 | homeobox protein (HOX-11) | 1209-1552 |  |
| D4j | M76541 | DNA-binding protein (NF-E1) | 706-1053 |  |
| D4ik | M76766 | transcription factor (TFIIB) | 407.769 |  |
| D41 | M80627 | HEB helix-loop-helix protein (HEB) | 3676-3984 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| D4m | M81601 | transcription elongation factor (SII) | 227-593 |  |
| A21 | M81750 | myeloid cell nuclear differentiation antigen | 549-873 |  |
| A51 | M81757 | S19 ribosomal protein | 113-408 |  |
| D4n | M81840 | NRL gene product | 946-1158 |  |
| D5a | M83234 | nuclease-sensitive element DNA-binding protein | 790-1099 |  |
| C2k | M84820 | retinoid X receptor beta (RXR-beta) | 643-1135 |  |
| C6g | M87338 | replication factor C 40-kDa subunit (A1) | 882-1286 |  |
| C6h | M87339 | replication factor C $37-\mathrm{kD}$ a subunit | 98-355 |  |
| D5b | M87503 | IFN-responsive transcription factor subunit | 1057-1520 |  |
| D5c | M92299 | homeobox 21 protein (HOX2A) | 1718-1945 |  |
| D5d | M92843 | zinc finger transcriptional regulator | 892-1271 |  |
| D5e | M93255 | FLI-1 | 728-1118 |  |
| E4e | M95489 | follicle stimulating hormone receptor | 1507-1752 |  |
| D5f | M96824 | nucleobindin precursor | 701-1068 |  |
| D5g | M96944 | B-cell specific transcription factor (BSAP) | 2446-2771 |  |
| D5h | M97287 | MAR/SAR DNA binding protein (SATB1) | 1921-2226 |  |
| D5i | M97676 | (region 7) homeobox protein (HOX7) | 1091-1450 |  |
| E4h | S64045 | 5HT1a=5-hydroxytryptamine receptor \{transmembrane regions 5 and 6\} | 128-413 |  |
| A5m | U01160 | transmembrane 4 superfamily protein (SAS) | 98-409 |  |
| B4g | U02081 | guanine nucleotide regulatory protein (NET1) | 1079-1323 |  |
| B4h | U02082 | guanine nucleotide regulatory protein (tim1) | 1852-2185 |  |
| D5j | U02326 | clone ndf43 neu differentiation factor | 1430-1701 |  |
| D5k | U02368 | PAX3/forkhead transcription factor fusion | 2231-2569 |  |
| D51 | U02619 | TFIIIC Box B-binding subunit | 5023-5369 |  |
| D5m | U02683 | alpha palindromic binding protein | 1630-2062 |  |
| A2m | V03056 | tumor suppressor (LUCA-1) | 2039-2444 |  |
| D5n | U03494 | transcription factor LSF | 1358-1681 |  |
| B4i | U03688 | dioxin-inducible cytochrome P450 (CYP1B1) | 1212-1556 |  |
| D6a | U04847 | Ini1 | 125-538 |  |
| D6b | U05040 | FUSE binding protein | 1002-1339 |  |
| $\overline{\text { A5n }}$ | $\cup 05340$ | p55CDC | 1236-1522 |  |
| B4i | U05875 | clone pSK1 interferon gamma receptor accessory factor-1 (AF-1) | 1702-2039 |  |
| B1c | U07139 | voltage-gated calcium channel beta subunit | 2008-2383 |  |
| B4k | U07236 | mutant lymphocyte-specific protein tyrosine kinase (LCK) | 930-1207 |  |
| Ā6a | U07616 | amphiphysin | 1740-2143 |  |
| B41 | U07707 | epidermal growth factor receptor substrate (eps15) | 1828-2140 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| Ē69 | U07819 | contactin 1 precursor (CNTN1) | 2735-3130 |  |
| D6c | U08015 | NF-ATC | 2039-2374 |  |
| D6d | U08191 | R kappa B | 4657-4920 |  |
| D6e | 008853 | transcription factor LCR-F1 | 1575-1928 |  |
| B4m | U09564 | serine kinase | 487-833 |  |
| D6f | U09579 | melanoma differentiation associated (mda-6) | 1745-2063 |  |
| B4n | U09607 | JAK family protein tyrosine kinase JAK3 | 3556-3892 |  |
| D6g | U10323 | nuclear factor NF45 | 967-1380 |  |
| D6h | U10324 | nuclear factor NF90 | 2901-3146 |  |
| D6i | U10421 | HOX A1 homeodomain protein (HOXA1) | 132-492 |  |
| D6j | U12535 | epidermal growth factor receptor kinase substrate (Eps8) | 2293-2645 |  |
| C21 | $\cup 13021$ | positive regulator of programmed cell death ICH-1L (lch-1) | $851-1218$ |  |
| D6k | U13897 | homolog of Drosophila discs large protein isoform 1 (hdig-1) | 2248-2624 |  |
| D61 | U14575 | (ard-1) | 665-942 |  |
| D6m | U14755 | LIM domain transcription factor LIM-1 (hLIM-1) | 479-759 |  |
| D6n | $U 15979$ | (dik) | 1090-1403 |  |
| B5a | U16031 | transcription factor IL-4 stat | 1816-2118 |  |
| C6i | X06745 | DNA polymerase alpha-subunit | 3721-4093 |  |
| A2n | $\times 07024$ | $X$ chromsome CCG1 protein inv in cell proliferation | 4002-4343 |  |
| A3a | $\times 15218$ | ski oncogene | 2354-2662 |  |
| A3b | X15219 | sno oncogene sno N protein ski-related | 2224-2652 |  |
| E6h | $\times 16841$ | N -CAM (a nontransmembrane isoform) from skeletal muscle | 2338-2646 |  |
| A3c | $\times 51630$ | Wilms tumor WT1 zinc finger protein Krueppel-like | 1866-2254 |  |
| D7a | X55122 | GATA-3 transcription factor | 1097-1383 |  |
| A6b | X55504 | P120 antigen | 1970-2245 |  |
| D7b | $\times 59738$ | ZFX put transcription activator isoform 1 | 749-1113 |  |
| D7c | $\times 67951$ | proliferation-associated gene (pag) | 543-856 |  |
| B5b | $\times 70326$ | MacMarcks | 638-1008 |  |
| B5c | X74979 | TRK E | 2138-2411 |  |
| E6i | Z26317 | desmoglein 2 | 2819-3135 |  |
| F7c | A00914 | angiotensin-converting enzyme (ACE) | 2123-2483 |  |
| F7d | A06925 | relaxin H2 | 123-427 |  |
| F7e | D10232 | renin-binding protein | 289-589 |  |
| E4i | D28538 | glutamate receptor type 1 subtype 5a | 3745-4027 |  |
| F7 | J04040 | glucagon | 201-540 |  |
| E4j | L19058 | glutamate receptor 5 | 2514-2779 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| F7g | M13981 | inhibin A-subunit | 828-1183 |  |
| F7h | M14200 | diazepam binding inhibitor | 67-257 |  |
| E4k | M15169 | Beta-2-adrenergic receptor | 2412-2783 |  |
| E41 | M29066 | dopamine d2 receptor | 1226-1521 |  |
| F7i | M31159 | growth hormone-dependent insulin-like growth factor-binding protein | 451-744 |  |
| F7j | M68867 | retinoic acid-binding protein II | 489-863 |  |
| E4m | M76446 | alpha A1 adrenergic receptor | 1599-1942 |  |
| E4n | M86841 | serotonin receptor type 2 | 938-1239 |  |
| F7k | U06863 | follistatin-related protein precursor | 1093-1425 |  |
| F71 | $\times 58022$ | corticotropin-releasing factor-binding protein | 853-1140 |  |
| AGC | HT0121 | cyclin-dependent kinase 2 | 1774-2180 |  |
| A6d | HT0191 | cell division cycle protein 25A tyrosine phosphatase | 1632-1978 |  |
| A6e | HT0285 | cyclin D3 | 537-894 |  |
| C6j | HT330 | single-stranded DNA-binding protein pur-alpha | 563-855 |  |
| A6! | HT0609 | cyclin A | 876-1218 |  |
| C6k | HT767 | DNA topoisomerase I | 2388-2796 |  |
| C61 | HT784 | DNA topoisomerase II alpha | 2459-2883 |  |
| C6m | HT1104 | 6-O-methylguanine-DNA methyltransferase | 241-546 |  |
| C6n | HT1175 | DNA excision repair protein ERCC2 $5^{\prime}$ end | 1520-1821 |  |
| A3d | HT1426 | prohibitin | 172-455 |  |
| A3e | HT1436 | proto-oncogene raf | 1704-1989 |  |
| C2m | HT1483 | glutathione reductase | 719-1057 |  |
| A3f | HT1489 | proto-oncogene c-abl tyrosine protein kinase alt transcript 1 | 3240-3612 |  |
| A6g | HT1547 | cyclin D1 | 3427-3784 |  |
| C2n | HT1790 | glutathione S-transierase 12 | 72-420 |  |
| C7a | HT1848 | DNA excision repair protein ERCC1 alt transcript 1 | 625-938 |  |
| C3a | HT2041 | glutathione S-transferase M1 | 504-906 |  |
| C3b | HT2042 | glutathione S-transferase pi | 203-511 |  |
| C3c | HT2168 | glutathione S-transferase A1 | 257-583 |  |
| A6h | HT2181 | cyclin D2 | 3932-4284 |  |
| A3g | HT2291 | proto-oncogene c-src1 tyrosine kinase domain | 893-1189 |  |
| A3h | HT2788 | proto-oncogene rel | 1357-1605 |  |
| A3i | HT2856 | proto-oncogene rhoA multidrug resistance protein | 290-572 |  |
| C3d | HT2859 | glutathione peroxidase | 454-745 |  |
| A3j | HT3039 | proto-oncogene shb src-2 homolog | 1365-1657 |  |
| C3e | HT3190 | apoptosis regulator bcl-x | 412-676 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| C7b | HT3218 | superoxide dismutase 1 cytosolic | 198-486 |  |
| C7c | HT3337 | DNA mismatch repair protein hmlh1 | 1765-2020 |  |
| A $\overline{\text { bi }}$ | HT3410 | cell division cycle protein 25 nucleotide exchange factor | 3372-3651 |  |
| A3k | HT3563 | tumor suppressor DCC colorectal | 3749-4042 |  |
| C̄3' | HT3614 | cytochrome P450 reductase | 789-1082 |  |
| C7d | HT4209 | xeroderma pigmentosum group $C$ repair complementing protein p58/HHR23B | 582-885 |  |
| C7e | HT4247 | xeroderma pigmentosum group C repair complementing protein HHR23A | 355-632 |  |
| A6j | HT4540 | cyclin H | 717-1026 |  |
| C3g | HT4547 | glutathione S-transterase T1 | 617-914 |  |
| C3h | HT5168 | ionizing radiation resistance-conferring protein | 856-1114 |  |
| E6j | J02703 | endothelial membrane glycoprotein IIIA (GPIIIA) | 2038-2373 |  |
| E6k | J04145 | neutrophil adherence receptor alpha-M subunit | 2888-3183 |  |
| E61 | J05633 | integrin beta- 5 subunit | 2279-2528 |  |
| E6m | L12002 | integrin alpha 4 subunit | 2709-3063 |  |
| E6n | M15395 | leukocyte adhesion protein (LFA-1/MAC-1/P15095 family) beta subunit | 2367-2664 |  |
| E7a | M34480 | platelet glycoprotein IIB (GPIIB) | 268-639 |  |
| E7b | M35198, J05522 | integrin B-6 | 1619-1901 |  |
| E7c | M59911 | integrin alpha-3 chain | 2562-2944 |  |
| E7d | M81695, Y00093 | leukocyte adhesion glycoprotein P15095 | 88-271 |  |
| E7e | X06256 | fibronectin receptor alpha subunit | 2094-2367 |  |
| E7f | X07979 | fibronectin receptor beta subunit | 2116-2482 |  |
| E7g | X53586 | integrin alpha 6 | 3642-3988 |  |
| E7h | X53587 | integrin beta 4 | 5357-5697 |  |
| E7i | X68742 | integrin alpha subunit | 2690-2976 |  |
| E7j | X74295 | alpha 7B integrin | 255-591 |  |
| E7k | Y00796 | leukocyte-associated molecule-1 alpha subunit (LFA-1 alpha subunit) | 4526-4856 |  |
| C3i | D38122 | Fas ligand | 516-840 |  |
| B7i | D49547 | heat-shock protein 40 | 1400-1782 |  |
| D7d | J03133 | transcription factor SP1 3' end | 1876-2272 |  |
| $\overline{\text { B }}$ d | L07032 | protein kinase C theta (PKC) | 2306-2601 |  |
| B5e | L26318 | protein kinase (JNK1) | 952-1263 |  |
| A 6 k | L27211 | CDK4-inhibitor (p16-INK4) | 482.836 |  |
| $\overline{\text { B }}$ f | L35253 | P38 mitogen activated protein (MAP) kinase | 925-1204 |  |
| 859 | L36719 | MAP kinase kinase 3 (MKK3) | 790-1169 |  |
| B5' | L36870 | MAP kinase kinase 4 (MKK4) | 2788-3103 |  |



| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| B6c | $\times 54637$ | tyk2 non-receptor protein tyrosine kinase | 3787-4110 |  |
| A4h | X56681 | jund | 508-780 |  |
| A4i | X59932 | c-src-kinase | 488-876 |  |
| B6d | X60188 | ERK1 protein serine/threonine kinase | 754-1094 |  |
| B6e | X80692 | ERK3 | 806-1267 |  |
| C31 | X86779 | FAST kinase | 865-1239 |  |
| E7m | X87838 | beta-catenin | 2061-2463 |  |
| C3m | X89986 | NBK apoptotic inducer protein | 935-1200 |  |
| A7d | X92669 | p35 cyclin-like CAK1-associated protein | 39-237 |  |
| B6f | Z29090 | phosphatidylinositol 3-kinase | 3021-3283 |  |
| C3n | L11015 | lymphotoxin-beta | 69-429 |  |
| B6g | L31951 | protein kinase (JNK2) | 638-1000 |  |
| B6h | L34583 | tyrosine phosphatase (clone HFAP10) | 1372-1701 |  |
| C4a | L41690 | TNF receptor-1 associated protein (TRADD) | 1009-1313 |  |
| C4b | M14745 | bcl-2 | 5087-5382 |  |
| C4c | U15172 | NIP1 (NIP1) | 412-719 |  |
| C4d | U15174 | NIP3 (NIP3) | 272-637 |  |
| C4e | U20537 | cysteine protease MCH2 isom beta (MCH2) | 387-697 |  |
| C4f | U23765 | BAK protein | 1371-1661 |  |
| C 4 g | U28014 | cysteine protease (ICEREL-II) | 763-1107 |  |
| C4h | U29680 | A1 protein | 64-293 |  |
| B6i | U34819 | JNK3 alpha2 protein kinase (JNK3A2) | 1018-1413 |  |
| C4i | U45878 | inhibitor of apoptosis protein 1 | 1444-1848 |  |
| C4j | U45879 | inhibitor of apoptosis protein 2 | 2000-2363 |  |
| C4k | U45880 | $X$-linked inhibitor of apotosis protein XIAP | 266-621 |  |
| C41 | U56390 | cysteine protease ICE-LAP6 | 986-1289 |  |
| C4m | U57059 | Apo-2 ligand | 211-616 |  |
| C4n | U60519 | apoptotic cysteine protease Mch4 (Mch4) | 2276-2690 |  |
| C5a | U60520 | apoptotic cysteine protease Mch5 isom alpha (Mch5) | 1327-1607 |  |
| B6j | X14454 | interferon regulatory factor 1 | 478-695 |  |
| C5b | X96586 | FAN protein | 2449-2726 |  |
| C5c | Y09392 | WSL-LR WSL-S1 and WSL-S2 proteins | 1407-1671 |  |
| D7h | D11117 | homeobox HOX 4A homeodomain protein | 4200-4447 |  |
| A7e | D38305 | Tob | 626-926 |  |
| B6k | D42108 | phospholipase C | 1635-2003 |  |
| D7i | D45132 | zinc-finger DNA-binding protein | 5113-5551 |  |


| Array Coordinate | GeneBank \# | Gene Name | Position |  |
| :---: | :---: | :---: | :---: | :---: |
| Ē5a | D49394 | serotonin 5-HT3 receptor | 1703-2000 |  |
| A4j | L16464 | ETS oncogene (PEP1) | 418-711 |  |
| A71 | L29216 | CLK2 | 1106-1356 |  |
| A7g | L29220 | CLK3 | 551-1002 |  |
| A7h | L29222 | CLK1 | 144-459 |  |
| E5b | L76224 | NMDA receptor | 2097-2395 |  |
| B7m | M11717 | heat shock protein (HSP 70) | 1962-2225 |  |
| F5g | M27544 | insulin-like growth factor | 652.919 |  |
| B61 | M68516 | protein C inhibitor | 8035-8423 |  |
| F5h | M86528 | neurotrophin-4 (NT-4) | 721-1079 |  |
| 86m | U09578 | MAPKAP kinase (3pK) | 486-837 |  |
| ATi | U10564 | CDK tyrosine 15-kinase WEE1HU (WEE1HU) | 1259-1502 |  |
| C7i | U12134 | DNA damage repair and recombination protein RAD52 | 1528-1733 |  |
| B6n | U14187 | receptor tyrosine kinase ligand LERK-3 (EPLG3) | 175-566 |  |
| B7a | U14188 | receptor tyrosine kinase LERK-4 (EPLG4) | 169-436 |  |
| B7b | U18087 | 3'5'-cAMP phosphodiesterase HPDE4A6 | 1119-1453 |  |
| C5d | U21092 | CD40 receptor associated factor 1 (CRAF1) | 980-1322 |  |
| A7j | U22398 | CDK-inhibitor P57KIP2 (KIP2) | 1048-1316 |  |
| A 4 k | U24166 | EB1 | 488-796 |  |
| A41 | U26710 | CBL-B | 3054-3444 |  |
| D7j | U28838 | transcription factor TFIIIB 90 kDa subunit (HTFIIIB90) | 2336-2605 |  |
| D7k | M30504 | transcription initiation factor TFIID subunit TAFII31 | 260-638 |  |
| F6n | U32659 | IL-17 | 257.578 |  |
| C5e | U32944 | cytoplasmic dynein light chain 1 (hdlc1) | 48-265 |  |
| B7c | U33635 | colon carcinoma kinase-4 (CCK4) | 3507-3784 |  |
| C7j | U33841 | ataxia telangiectasia (ATM) | 8938-9135 |  |
| A7k | U35735 | RACH1 (RACH1) | 1072-1391 |  |
| C5f | U39613 | cysteine protease ICE-LAP3 | 541-844 |  |
| B7d | U39657 | MAP kinase kinase 6 (MKK6) | 1060-1389 |  |
| B7e | U40282 | integrin-linked kinase (ILK) | 1245-1530 |  |
| A71 | U41816 | C-1 | 143-356 |  |
| D71 | U43188 | Ets transcription factor (NERF-2) | 1967-2400 |  |
| B71 | U43408 | tyrosine kinase (Tnk1) | 1455-1849 |  |
| A4m | U57456 | transforming growth factor-beta signaling protein-1 (bsp-1) | 1417-1679 |  |
| C5g | U59747 | Bcl-w (bcl-w) | $121-403$ |  |
| D7m | U59863 | TRAF-interacting protein 1-TRAF | 674-887 |  |



## Mouse Array

In the mouse array according to the subject invention, all of the unique polynucleotide probe compositions will correspond to a mouse gene of interest. Mouse genes that are represented on the array are key genes, by which is meant that they have been reported to play primary roles in a variety of different biological processes. Typically the mouse genes represented on the array are genes that are under tight transcriptional control. Genes of interest that may be represented on the array include: oncogenes, cell cycle genes, apoptosis genes, growth factor genes, cytokine genes, interleukin genes, receptor genes, and genes associated with different stages of embryonic development.

In certain embodiments, of particular interest is an array having the following types of genes represented on its surface: oncogenes \& tumor suppressors; cell cycle regulators; stress response proteins; ion channel \& transport proteins; intracellular signal transduction modulators \& effectors; apoptosis-related proteins; DNA synthesis, repair \& recombination proteins; transcription factors \& general DNA binding proteins; growth factor \& chemokine receptors; interleukin \& interferon receptors, hormone receptors; neurotransmitter receptors; cell-surface antigens \& cell adhesion proteins; interleukins \& interferons; cytoskeleton \& motility proteins; and protein turnover. In a specific mouse array of interest, the spots are as listed in Table 2.

The mouse array of the subject invention finds use in a variety of different applications, where such applications include: profiling differential gene expression in transgenic knockout mice or other experimental mouse models; investigating processes such as embryo genesis and tumorigenesis; discovering potential therapeutic and diagnostic drug targets; and the like.

| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| D13473 | MmRad51; yeast DNA repair protein Rad51 and E coli RecA homologue. | C6m | 855-1199 |
| D17630 | Interleukin-8 receptor | E3h | 664-1022 |
| D25281 | Catenin alpha | E5m | 1276-1594 |
| D31788 | BST-1; lymphocyte differentiation antigen CD38 | B2h | 674-1014 |
| D31942 | Oncostatin M | F3n | 1017-1360 |
| L05630 | C5A receptor | E1g | 841-1165 |
| L07264 | Heparin-binding EGF-like growth factor (Diphtheria toxin receptor) | F2d | 258.673 |
| $\bigcirc 04807$ | Fms-related tyrosine kinase 3 Fli3/Flk2 ligand | C3i | 46-418 |
| L24495 | CD27; lymphocyte-specific NGF receptor family member | C21 | 596-846 |
| M28998 | Fibroblast growth factor receptor Basic (b FGF-R) | E2c | 200-583 |
| M58288 | Granulocyte colony - stimulatings factor receptor | E1i | 251-529 |
| M62301 | Growth/ diffterentiation factor 1 (GDF-1) (TGF- beta family) | F2b | 2267-2566 |
| M69042 | PKC-delta; protein kinase C delta type | B69 | 1740-2011 |
| M74517 | GA binding protein beta-2 chain | D3d | 613-931 |
| M83312 | CD 40L receptor (TNF receptor family) | E1f | 417-754 |
| M83649 | Fasl receptor (Fas antigen, Apo-1 antigen) | C3f | 416-736 |
| M86671 | Interleukin 12 (p40) beta chain | F4n | 652-963 |
| M95200 | Vascular endothelial growth factor (VEGF) | F4j | 688-955 |
| ŪO3421 | Interleukin 11 (adipogenesis inhibitory factor) | F4m | 196-475 |
| U14332 | Interleukin 15 | F5a | 605-1057 |
| U15159 | LIMK; LIM serine/threonine kinase | B51 | 1376-1699 |
| U83628 | DAD-1; defender against cell death 1 | C3d | 221-509 |
| U25416 | CD 30L receptor ( Lymphocyte activation antigene CD 30, Ki -1 antigene) | C2m | 135.435 |
| U44725 | Mast cell factor | F3i | 79-417 |
| U56819 | C-C chemokine receptor (Monocyte chemoattractant protein 1 receptor (MCP-1RA) | E1d | 965-1262 |
| X06381 | Leukemia inhibitory factor (LIF) (cholinergic differentiation factor) | F3d | 63-366 |
| X52264 | Intercellular adhesion molecule-1 | E7i | 1053-1385 |
| X59769 | Interleukin-1 receptor type II | E2n | 883-1134 |
| $\times 72305$ | Corticotropin releasing factor receptor | E1h | 1411-1748 |
| X72307 | Hepatocyte growth factor (hepapoitein) | F2e | 641.965 |
| 222703 | Keratinocyte growth factor FGF. 7 | F3b | 63-325 |
| 231663 | Activin type I receptor | E1a | 847-1130 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| D01034 | Transcription factor TF II D | B4j | 291-556 |
| D14340 | ZO-1; Tight junction protein; discs-large family member, partially homologous to a dlg-A tumor suppressor in Drosophilal | A2d | 3714-4001 |
| D16306 | ERCC5 excision repair protein; DNA-repair protein complementing XP-G cells (XPG) | C6i | 1336-1639 |
| L22472 | Bax; $\mathrm{Bcl}-2$ heterodimerization partner and homologue | C1g | 172-534 |
| L25606 | B7-2; T lymphocyte activation antigen CD86; CD28 antigen ligand 2, B7-2 antigen; alternative CTLA4 counter-receptor | B2g | 570-967 |
| L27105 | NF2; Merlin (moesin-ezrin-radixin-like protein); shwannomin, murine neurofibromatosis type 2 susceptibility protein | A1i | 2175-2400 |
| M13945 | Pim-1 proto-oncogene | A4a | 2713-2930 |
| M20157 | Egr-1 Zn -finger regulatory protein | D2i | 399-753 |
| M25811 | PKC-alpha; protein kinase C alpha type | B6e | 1566-1924 |
| M27129 | CD44 antigen | E6e | 789-1141 |
| M31042 | T-lymphocyte activated protein | D6h | 285-606 |
| M31131 | Neuronal-cadherin (N-cadherin) | E7k | 1212-1409 |
| M38700 | ATP-dependent DNA helicase II 70 kDa subunit; thyroid Ku (p70/p80) autoantigen p70 subunit; p70 Ku) | C5h | 274-632 |
| M63660 | G13; G-alpha-13 guanine nucleotide regulatory protein | B6n | 2057-2377 |
| M83380 | Transcription factor RelB | D7c | 1456-1728 |
| M84487 | Vascular cell adhesion protein 1 | E7m | 984-1304 |
| S71186 | ERCC3 DNA repair helicase; DNA-repair protein complementing XP-B cells (XPBC) | C6e | 1147-1444 |
| S76657 | CRE-BP1; cAMP response element binding protein 1 | B31 | 412.748 |
| U02887 | XRCC1 DNA-repair protein, affecting ligation | C7n | 900-1183 |
| U53228 | Nuclear hormone receptor ROR-ALPHA-1 | D5i | 368-675 |
| U57311 | 14-3-3 protein eta | B79 | 374-640 |
| X56135 | Prothymosin alpha | A7m | 186-455 |
| X57487 | PAX-8 (paired box protein PAX 8) | D51 | 680-1011 |
| X58995 | CamK IV; Ca2/calmodulin-dependent protein kinase IV (catalytic chain) | B5! | 1269-1608 |
| $\times 66323$ | ATP-dependent DNA helicase II 80 kDa subunit; thyroid Ku (p70/p80) autoantigen $p 80$ subunit; $p 80 \mathrm{Ku}$ ) | C5i | 565-875 |
| $\begin{aligned} & \times 67812 \\ & \times 68193 \end{aligned}$ | Ret proto-oncogene (Papillary thyroid carcinoma-encoded protein) Nm23-M2; nucleoside diphosphate kinase B; metastasis-reducing protein; c-myc-related transcription factor | A4i | $2359-2680$ $80-454$ |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| X97052 | MAPKK6; MAP kinase kinase 6(dual specificity) (MKK6) | B6d | 375-711 |
| D17384 | DNA polymerase alpha catalytic subunit (p180) | C5I | 563-908 |
| D28492 | Caspase-3; Nedd2 cysteine protease (positive regulator of programmed cell death $\mathrm{ICH}-1$ homologue) | C1b | 398-694 |
| D50621 | PSD.95/SAP90A | D6d | 1512-1889 |
| J04946 | Angiotensin-converting enzyme (ACE) (clone ACE.5.) | F6f | 850-1113 |
| L08235 | Clusterin; complement lysis inhibitor; testosterone-repressed prostate message 2; apolipoprotein J; sulfated glycoprotein-2 | C3b | 515-744 |
| L12721 | Adipocyte differentiation-associated protein | D1c | 404-709 |
| L21671 | Epidermal growth factor receptor kinase substrate EPS8 | D2k | 1592-1873 |
| L33768 | Jak3 tyrosine-protein kinase; Janus kinase 3 | B5j | 3123-3426 |
| L33779 | Desmocollin 2 | E61 | 1317-1691 |
| L47650 | Stat6; signal transducer and activator of transcription 6; IL-4 Stat; STA6 | B4g | 2057-2411 |
| M12056 | Lymphocyte-specific tyrosine-protein kinase LCK | A5a | 1205-1488 |
| M22115 | ERA-1 Protein (ERA-1-993) | D21 | 723-1062 |
| M26283 | Homeo Box protein 2.1 (Hox-2.1) | D4a | 647-884 |
| M32309 | Zinc finger X-chromosomal protein (ZFX) | D7n | 2153-2554 |
| M55512 | WT1; Wilms tumor protein; tumor suppressor | A2c | 1262-1563 |
| M57422 | Tristetraproline | B4k | 262-504 |
| M96823 | Nucleobindin | D5j | 80-357 |
| M97013 | PAX-5 (B cell specific transcription factor) | D6a | 286-629 |
| S69336 | IFNgR2; interferon-gamma receptor second (beta) chain; interferon gamma receptor accessory factor-1 (AF-1) | B3b | 832-1089 |
| 574227 | Transcriptional enhancer factor 1 (TEF-1) | D7i | 934-1233 |
| U02079 | Transcription factor NFAT 1, isoform alpha | D7a | 1601-1910 |
| U05252 | DNA-binding protein SATB1 | D2e | 1101-1380 |
| U20372 | CCHB3; calcium channel (voltage-gated; dihydropyridine-sensitive; L-type) beta-3 subunit) | B2c | 351-639 |
| U20553 | p57kip2; cdk-inhibitor kip2 (cyclin-dependent kinase inhibitor 1B) member of the p21CIP1 Cdk inhibitor family; candidate tumor suppressor gene | A7g | 989-1272 |
| U36203 | snoN; ski-related oncogene | E2j | 671-1006 |
| $\times 14759$ | Homeo Box protein 7.1 (Hox-7.1) | D4f | 740-992 |
| $\times 14943$ | Neuronal cell surface protein F3 | E71 | 1033-1311 |
| $\times 55123$ | GATA-3 transcription factor | D3f | 858-1125 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| X 57621 | YB1 DNA binding protein | D7j | 550-873 |
| X 58384 | Dipeptidyl peptidase iv | E7f | 61-294 |
| X 59421 | Fli-1 ets-related proto-oncogene | A3b | 267-623 |
| X 66224 | RXR-beta cis-11-retinoic acid receptor | B4c | 1225-1477 |
| X78445 | C3H cytochrome P450; Cyp1b1 | B1j | 295-593 |
| Х 96859 | Ubiquitin-conjugating enzyme, yeast Rad6 homologue; murine HR6B | C7k | 51-392 |
| 227088 | Relaxin | C4i | 51-365 |
| 227410 | Transcription factor LIM-1 | D6m | 1673-1934 |
| D10061 | DNA topoisomerase I (Top I) | C5m | 1051-1357 |
| D12513 | DNA topoisomerase II (Top II) | C5n | 520-870 |
| D30687 | GST Pi 1; glutathione S-transferase Pi 1; preadipocyte growth factor | C2d | 62-369 |
| J03958 | Glutathione S-transferase A | C1n | 54-311 |
| J04696 | Glutathione S-transferase Mu 1 | C2b | 13-263 |
| L10656 | c-Abl proto-oncogene | A4k | 878-1145 |
| M13071 | A-Raf proto-oncogene | A3k | 1042-1320 |
| M17031 | c-Src proto-oncogene | A4n | 452-758 |
| M35523 | Retinoic acid binding protein II cellular (CRABP-II) | D6e | 276-571 |
| M83749 | Cyclin D2 (G1/S-specific) | A6g | 781-1074 |
| U43844 | Cyclin D3 (G1/S-specific) | A6h | 484-790 |
| S49542 | 5-Hydroxytryptamine receptor [Serotonin receptor type 2 (5HT2)] | E4e | 400-707 |
| S78355 | Cyclin D1 (G1/S-specific) | A6i | 1858-2205 |
| U02098 | Pur-alpha transcriptional activator; sequence-specific ssDNA-binding protein | C7e | 1082-1309 |
| U27323 | Cdc25a; cdc25M1; MPI1 (M-phase inducer phosphatase 1) | A7j | 606-986 |
| $\times 07414$ | ERCC-1: DNA excision repair protein | C6d | 189-484 |
| X15842 | c-rel proto-oncogene | A2m | 1729-2064 |
| X69618 | inhibin alpha subunit | F2g | 810-1117 |
| X76341 | Glutathione reductase | C1m | 115-377 |
| X81581 | Insulin-like growth factor binding protein-3 (IGFBP-3) | F2k | 474-719 |
| Z26580 | Cyclin A (G2/M-specific) | A6a | 701-1009 |
| 246845 | Preproglucagon | A5i | 172-531 |
| M61909 | NF-kB p65; NF-kappa-B transcription factor p65 subunit; rel-related polypeptide | B4a | 101-363 |
| D11091 | PKC-theta; protein kinase C theta type | B6h | 658-957 |
| D13867 | VLA-3 alpha subunit | E7n | 288-589 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| D17571 | NADPH-cytochrome P450 reductase | C4a | 326-605 |
| D17584 | Beta-protachykinin a | A5j | 273-523 |
| D30743 | Wee 1/p87; cdc2 tyrosine 15-kinase | A7h | 1816-2159 |
| D83966 | Protein tyrosine phosphatase | C4g | 1060-1429 |
| J05205 | Jun-D; c-jun-related transcription factor | A3g | 737-964 |
| L23423 | Integrin alpha 7 | E7e | 2399-2713 |
| L28177 | Gadd45; growth arrest and DNA-damage-inducible protein | C3j | 144-434 |
| L35049 | Bcl-xL apoptosis regulator (bcl-x long); Bcl-2 family member | C1j | 641-906 |
| X03919 | N -myc proto-oncogene protein | A3j | 3262-3450 |
| M20473 | cAMP-dependent protein kinase type I-beta regulatory chain | B5g | 538.750 |
| M21065 | IRF1; interferon regulatory factor 1 | B7k | 1-233 |
| M36830 | HSP86; heat shock 86kD protein | B1d | 255-551 |
| M60778 | LFA1-alpha; integrin alpha L; leukocyte adhesion glycoprotein LFA-1 alpha chain; antigen CD11A (p180) | B3e | 1838-2050 |
| M88127 | APC; Adenomatous Polyposis Coli protein | A1a | 4127-4476 |
| S93521 | Cdc25b; cdc25M2; MPI2 (M-phase inducer phosphatase 2) | A7k | 1893-2200 |
| U03279 | P13-K p110; phosphatidylinositol 3-kinase catalytic subunit | B6j | 1437-1723 |
| U03560 | HSP27; heat shock 27kD protein 1 | B1a | 245-500 |
| U05247 | Csk; c-Src-kinase and negative regulator | B4n | 645-984 |
| U06948 | Fasl; Fas antigen ligand; generalized lymphoproliferation disease gene (gld) in mice | C3g | 168-488 |
| U10871 | MAPK; MAP kinase; p38 | B5m | 465-780 |
| U19597 | p19ink4; cdk4 and cdk6 inhibitor | A7d | 228-516 |
| U19617 | Ell-1 Ets family transcription factor | D2j | 1585-1902 |
| U21050 | CRAF1; TNF receptor (CD40 receptor) associated factor; TRAF-related | C3c | 1225-1466 |
| U25844 | SPI3; serpin; similar to human proteinase inhibitor 6 (placental thrombin inhibitor) serine proteinase inhibitor | C41 | 915-1230 |
| U25995 | RIP cell death protein; Fas/APO-1 (CD95) interactor, contains death domain | C4i | 1945-2223 |
| U29056 | SLAP; src-like adapter protein; Eck receptor tyrosine kinase-associated | B5c | 109-427 |
| U43678 | Alm; ataxia telangiectasia murine homologue | C5g | 8989-9170 |
| U51196 | EB1 APC-binding protein | A1e | 607-834 |
| U51907 | TANK; I-TRAF; TRAF family member associated NF-kB activator | B4h | 135-437 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| U59463 | Caspase-11; ICH-3 cysteine protease; upstream regulator of ICE | C1a | 352-686 |
| U59883 | MLH1 DNA mismatch repair protein; MutL homologue | C6k | 1037-1278 |
| X04480 | Insulin-like growth factor-IA | F3a | $183-406$ |
| $\times 07640$ | Cell surface glycoprotein MAC-1 alpha subunit | E6j | 1892-2179 |
| X13664 | N -ras proto-oncogene; transforming G-protein | A5e | 548-857 |
| X13945 | L-myc proto-oncogene protein | A3h | 5287-5590 |
| X14951 | CD18 antigen beta subunit (leukocyte adhesion LFA-1) (CD3, P150, 95) | E5n | 1366-1706 |
| $\times 52191$ | c-Fgr proto-oncogene | A4m | 1305-1538 |
| X53176 | Integrin alpha 4 | E7b | 2176-2449 |
| X53532 | PKC-beta; protein kinase C beta-II type | B6i | 1712-2089 |
| $\times 53584$ | HSP60; heat shock 60 kDa protein 1 (chaperonin, GroEL homologue); mitochondrial matrix protein P1 | B1b | 1432-1459 |
| X57111 | c-Cbl proto-oncogene (Adaptor protein) | A5b | 858-1151 |
| X59868 | Cdc25 phosphatase; guanine nucleotide releasing protein | A7i | 942-1276 |
| $\times 60671$ | Ezrin; Villin 2; NF-2 (merlin) related filament/plasma membrane associated protein | A11 | 1571-1812 |
| X64713 | Cyclin B1 (G2/M-specific) | A6c | 1184-1447 |
| X69902 | Integrin alpha 6 | E7d | $261 \cdot 611$ |
| X72395 | 5-Hydroxytryptamine (serotonin) receptor 3 | E4j | 1422-1711 |
| $\times 73573$ | Homeobox protein HOXD-3 | D4h | 141-362 |
| $\times 75888$ | Cyclin E (G1/S-specific) | A6i | 799-1140 |
| X76850 | MAPKAPK-2; MAP kinase-activated protein kinase; MAPKAP kinase 2 | B5n | 719.987 |
| X83971 | Fra-2 (fos-related antigen 2) | A3d | 617-844 |
| $\times 84311$ | Cyclin A1 (G2/M-specific) | A6b | 656.916 |
| $\times 85788$ | $\overline{\mathrm{DCC}}$; netrin receptor; immunoglobulin gene superfamily member; former tumor suppressor protein candidate | A1d | 4193-4508 |
| X92410 | MHR23A; Rad23 UV excision repair protein homologue; xeroderma pigmentosum group C (XPC) repair complementing protein | C6i | 613-955 |
| X92411 | MHR23B; Rad23 UV excision repair protein homologue; xeroderma pigmentosum group C (XPC) repair complementing protein | C6j | 542-807 |
| Y00769 | Integrin beta | E7g | 1990-2320 |
| Z32767 | MmRad52; yeast DNA repair protein Rad52 homologue | C6n | 159.417 |
| 237110 | Cyclin G (G2/M-specific) | A6k | 300-619 |
| D13458 | Prostaglandin E2 receptor EP4 sublype | B3I | 1146-1442 |
| D90205 | Interleukin-5 receptor | E3i | 1389-1739 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| J00380 | Epidermal growth factor (EGF) | F1j | 180-505 |
| J04843 | Erythropoietin receptor | E2a | 1193-1377 |
| J05149 | Insulin receptor | E4a | 653-1011 |
| K01700 | p53; tumor suppressor; DNA-binding protein | A11 | 1125-1517 |
| L03529 | Ci2r; coagulation factor II (thrombin) receptor | B2j | 762-1154 |
| L09562 | PTPRG; protein-tyrosine phosphatase gamma | B71 | 1248-1504 |
| L10075 | DNA-binding protein SMBP2 | D2f | 4790-5088 |
| L12120 | Interleukin-10 receptor | E3a | 1762-2110 |
| L20048 | Interleukin-2 receptor gamma chain | E3c | 1073-1313 |
| L24755 | Bone morphogenetic protein 1 | F1b | 2402-2676 |
| L33406 | Uromodulin | F4i | 1809-2136 |
| L34169 | Thrombopoietin | F4e | 652-954 |
| M13177 | Transforming growth factor beta | F4i | 772-1075 |
| M13926 | Granulocyte colony-stimulating factor (G-CSF) | F2a | 86-377 |
| M14220 | Neuroleukin | F3m | 1110-1490 |
| M14951 | Insulin-like growth factor-2 (somatomedin A) | F2n | 46-328 |
| M15131 | Interleukin 1 beta | F4k | 827-1225 |
| M16449 | c-myb proto-oncogene protein | A2k | 1212-1513 |
| M16819 | Tumor necrosis factor beta TNF-beta (Lymphotoxin-alpha) | F4h | 461-805 |
| M20658 | Interleukin-1 receptor | C3n | 2050-2410 |
| $\times 05010$ | CSF-1; M-CSF; colony stimulating factor-1 | A5g | 1268-1657 |
| M27959 | Interleukin-4 receptor (membrane-bound form) | E3e | 2469-2705 |
| M28233 | Interferon-gamma receptor | E2m | 1262-1550 |
| M29697 | Interleukin-7 receptor | E3g | 701-1104 |
| M34815 | Gamma interferon induced monokine (MIG) | F1m | 42-323 |
| M37897 | Interleukin 10 | F41 | 175-456 |
| M57999 | NF-kappa B binding subunit (nuclear factor) (TFDB5) | D5g | 3122-3417 |
| M59378 | Tumor necrosis factor receptor 1; TNFR-1 | C5d | 1961-2376 |
| M84607 | PDGFRa; platelet-derived growth factor alpha-receptor | A4e | 474-803 |
| M84746 | Interleukin-9 receptor | E3i | 795-1086 |
| M87039 | iNOSI; nitric oxide synthase (inducible) | C3m | 3178-3455 |
| M89641 | Interferon alpha-beta receptor | E21 | 808-1120 |
| M94087 | Activating transcription factor 4 (mATF4) | D1b | 416-769 |
| S56660 | Beta2-RAR; retinoic acid receptor beta-2 | B3k | 589-896 |
| S67051 | Tie-2 proto-oncogene | A4i | 1843-2179 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| U00182 | IGF-I-R alpha; insulin-like growth factor I receptor alpha subunit | C31 | 489-885 |
| U04710 | IGFR II: insulin-like growth factor receptor II, cation-independent mannose-6-P receptor; elevated in Wilms's tumor cells | C3k | 707-1060 |
| U06922 | Stat3; APRF; acute phase response factor | B4e | 1575-1910 |
| U18542 | Calcitonin receptor 1b | E3k | 1375-1630 |
| U32329 | Endothelin b receptor [Ednrb] | E1i | 279-695 |
| U32330 | Prepro-endothelin-3 | F4c | 703-1008 |
| X04367 | Pre-platelet-derived growth factor receptor | E2i | 2336-2677 |
| X04836 | CD 4 receptor (T cell activation antigene) | E1e | 1652-1877 |
| X07962 | Interleukin 7 | F5d | 241-496 |
| X12531 | Macrophage inflamatory protein | F3e | 25-359 |
| X14432 | Thrombomodulin | F4d | 1082-1365 |
| X51975 | Interleukin 6 (B cell differentiation factor) | F5C | 1638-1898 |
| X53779 | Androgen receptor | E3j | 2189-2491 |
| X56848 | Bone morphogenetic protein 4 (BMP-4) (TGF-beta family) | F1d | 1275-1513 |
| $\times 57349$ | Translerrin receptor protein (p90, CD71) | B3h | 654-1023 |
| X57413 | Transforming growth factor beta 2 | F4g | 2227-2541 |
| X57497 | Glutamate receptor, ionotropic AMPA 1 | E5h | 1290-1657 |
| X57796 | TNF 55; tumor necrosis factor 1 (55kd) | C5b | 656-1022 |
| X58876 | Mdm2; p53-regulating protein | A1h | 1364-1646 |
| X61753 | Transcription factor 1 for heat shock gene | D6i | 203-570 |
| X65453 | CD40L; CD40 ligand | C2n | 545-809 |
| $\times 68932$ | c-Fms proto-oncogene (macrophage colony stimulating factor 1 (CSF-1) receptor) | A4b | 2399-2686 |
| $\times 70472$ | $B-m y b$ proto-oncogene; myb-related protein $B$ | A2f | 2109-2456 |
| $\times 76654$ | Ear-2; v-erbA related proto-oncogene | A2n | 1065-1376 |
| $\times 80764$ | Tie-1 tyrosine-protein kinase receptor | B3g | 1425-1844 |
| D10651 | Glutamate receptor, ionotropic NMDA2B (epsilon 2) | E5j | 506-786 |
| D10217 | Glutamate receptor, ionotropic NMDA2A (epsilon 1) | E5i | 3966-4209 |
| D10329 | CD7 antigen | E6g | 28-421 |
| D00926 | Transcription factor S - 11 (transcription elongation factor) | D7d | 518-767 |
| D12482 | Basic Fibroblast growth factor (b- FGF) | F1a | $290 \cdot 620$ |
| D16250 | Bone morphogenetic protein receptor | E1c | 1454-1837 |
| D17292 | G-protein-coupled receptor | E2d | 833-1115 |
| D17407 | Transcription factor SP2 | D79 | 734-1079 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| D29678 | Cdk5; cyclin-dependent kinase 5 | A6n | 552-882 |
| D25540 | TGF-beta receptor type 1 | E2k | 1407-1629 |
| D26077 | Kinesin like protein KIF 3B | F6a | 3519-3722 |
| D29951 | Kinesin family protein KIF1A | F5m | 2553-2830 |
| D38258 | Fibroblast growth factor 9 | F1k | 91-379 |
| D83698 | Neuronal death protein | C4b | 627-805 |
| D84372 | Syp; SH-PTP2; adaptor protein tyrosine phosphatase | B5e | 1229-1543 |
| J03168 | Interferon regulatory factor 2 (IRF 2) | D41 | 718-976 |
| J02870 | Lamimin receptor 1 | E7j | 368-675 |
| D90176 | NF-1B protein (transcription factor) | D5f | 452-791 |
| J03236 | Jun-B; C-jun-related transcription factor | A3f | 514.740 |
| J03520 | Tissue plasminogen activator | F7e | 622-1020 |
| J 03770 | Homeo Box protein 4.2 (Hox-4.2) | D4e | 565-945 |
| J04113 | Nur77 early response protein; thyroid hormone (TR3) receptor | C4d | 825-1059 |
| J04103 | Ets-2 transcription factor | D3b | 917-1281 |
| J04115 | c-Jun proto-oncogene (transcription factor AP-1 component) | A2i | 951-1238 |
| J05609 | Serine protease inhibitor homolog J6 | F71 | 581-855 |
| K01759 | Nerve growth factor beta (beta-NGF) | F31 | 642-901 |
| L01640 | Cdk4; cyclin-dependent kinase 4 | A6m | 230-616 |
| K02582 | Acetylcholine receptor delta submit | E41 | 1400-1655 |
| L02526 | MAPKK1; MAP kinase kinase 3 (dual specificity) (MKK1) | B6a | 1284-1583 |
| L04662 | GABA-A transporter 4 | E5g | 960-1341 |
| L04663 | GABA-A transporter 3 | E5f | 1010-1320 |
| L07297 | Vegfr1; Vascular endothelial growth factor receptor $1 /$ Fms-related tyrosine kinase 1 (FII1) | A4j | 1144.1541 |
| L10084 | Adrenergic receptor, beta 1 | E4m | 404-772 |
| L25890 | Eph3 (Nuk) tyrosine-protein kinase receptor | B2k | 2255-2491 |
| L16953 | MTJ1; Dnaj-like heat-shock protein from mouse tumor | B1e | 1059-1384 |
| L19622 | TIMP-3 tissue inhibitor of metalloproteinases-3 | F7n | 274-592 |
| L24563 | Insulin receptor substrate-1 (IRS-1) | E4b | 1027-1304 |
| L13968 | YY1 (UCRBP) transcriptional factor | D7k | 1052-1292 |
| L28095 | Interleukin-converting enzyme (ICE) | F7a | 30-269 |
| L38847 | Hepatoma transmembrane kinase ligand | F2f | 927-1219 |
| L36179 | Voltage-gated sodium channel | B2f | 4179-4505 |
| L37296 | Bad; heterodimeric partner for $\mathrm{Bcl}-\mathrm{XL}$ and $\mathrm{Bcl}-2$; promotes cell death | C1d | 1079-1375 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| L35236 | Jnk stress-activated protein kinase (SAPK) | B5k | 795-1032 |
| M11686 | Cytoskeletal epidermal keratin (18 human) | F5i | 473.773 |
| M11434 | Nerve growth lactor alpha (alpha-NGF) | F3k | 294-494 |
| M10937 | Epidermal keratin (1 human) | F5k | 326-683 |
| M14537 | Nicotinic acetylcholine receptor | E5k | 1226-1568 |
| M14757 | $\bar{M}$ DR1; P-glycoprotein; multidrug resistance protein; efflux pump | B19 | 1500-1886 |
| M18934 | CD2 antigen | E6a | 354-602 |
| M17192 | Homeo Box protein 1.1 (Hox-1.1) | D3n | 466-723 |
| M19436 | Fetal myosin alkali light chain | F51 | 205-504 |
| M25892 | Interleukin 4 | F5b | 77-310 |
| M26391 | Rb; pp105; Retinoblastoma susceptibility-associated protein (tumor suppressor gene; cell cycle regulator) | A1m | 2036-2296 |
| M28489 | Rsk; ribosomal protein S6 kinase | B6i | 1191-1436 |
| M29464 | Pletelet- derived growth factor (A chain) (PDGF-A) | F4b | 152-425 |
| M28698 | Cytoskeletal epidermal keratin (19 human) | F5 | 194-500 |
| M29475 | RAG-1; V(D)J recombination activating protein | C7g | 2155-2404 |
| M29855 | Interleukin-3 receptor | E3d | 1975-2254 |
| M30642 | K-fibroblast growth factor | F3c | 309-577 |
| M34381 | Octamer binding transcription factor (Oct 3) | D5k | 774-999 |
| M33960 | Plasminogen activator inhibitor | F7h | 1096-1344 |
| M33158 | CD3 antigen, delta polypeptide | E6c | 73-361 |
| M34857 | Homeo Box protein 2.5 (Hox-2.5) | D4c | 11-277 |
| M36829 | HSP84; heat shock 84kD protein | B1c | 342-366 |
| M55617 | Mast cell protease (MMCP) - 4 | F7b | 634-992 |
| M61177 | Erk 1; exiracellular signal-regulated kinase 1; p44; Ert2 | B5h | 115-373 |
| M60651 | PI3-K p85; phosphatidylinositol 3-kinase regulatory subunit; phosphoprotein p85; PDGF signaling pathway member | B6k | 981-1260 |
| M58633 | p58/GTA; galactosyltransferase associated protein kinase (cdc2-related protein kinase) | A7b | 1022:1284 |
| M64086 | Serine protease inhibitor 2 (spi-2) | F7j | 1499-1754 |
| M64429 | B-Raf proto-oncogene | A31 | 1651-2036 |
| M68513 | Etk1 (Mek4; HEK) tyrosine-protein kinase receptor HEK | B21 | 2681-2915 |
| M64796 | RAG-2; V(D)J recombination activating protein | C7h | 671.944 |
| M84324 | Collagenase type IV | F6k | 696-1040 |
| M83336 | Interleukin-6 receptor beta chain; membrane glycoprotein gp130 | B3C | 1423-1741 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| M76601 | Alpha cardiac myosin heavy chain | F5e | 2094-2391 |
| M84819 | Retinoic acid receptor RXR-gamma | D6f | 701-1082 |
| M85078 | Granulocyte-macrophage colony-stimulating factor receptor | E2e | 904-1289 |
| M86566 | GABA-A receptor alpha-1 submit | E5d | 1251-1606 |
| M93428 | Endothelial ligand for L-selectin (GLYCAM 1) | F1i | 182-541 |
| M95633 | Integrin beta 7 subunit | E7h | 2142-2423 |
| U00478 | DNAse I | C6c | 665-871 |
| U03184 | Cortactin; protein tyrosine kinase substrate | B7h | 426-653 |
| U05672 | Adenosine A2M2 receptor | C2g | 491-735 |
| U04674 | DNA ligase I | C5j | 1678-2054 |
| 005671 | Adenosine A1M receptor | C2f | 302-673 |
| U04443 | Non-muscle myosin light chain 3 | F6b | 84-370 |
| U06119 | Cathepsin H | F6i | 325-694 |
| U06924 | Stat1; signal transducer and activator of transcription | B4d | 1749-2104 |
| U09507 | p21/Cip1/Waf1; cdk-inhibitor protein 1 | A7e | 9-403 |
| U11822 | Cdk7; MO15; cyclin-dependent kinase 7 (homologue of Xenopus MO15 cdk-activating kinase) | A7a | 454-824 |
| U10440 | p27kip1; G1 cyclin-Cdk protein kinase inhibitor, p21-related | A71 | 270-454 |
| U10551 | Gem; induced, immediate early protein; Ras family member | B7a | 220-471 |
| U12570 | VHL; Von Hippel-Lindau tumor suppressor protein | A2b | 885-1111 |
| U12983 | Cek 5 receptor protein tyrosine kinase ligand | F1g | 1037-1287 |
| U13705 | Glutathione peroxidase (plasma protein); selenoprotein. | C11 | 766-1046 |
| U14135 | Integrin alpha 5 (CD51) | E7c | 2170-2516 |
| U14173 | Ski proto-oncogene | A49 | 707-1037 |
| U17698 | Äblphilin-1 (abi-1) similar to HOXD3 | D1a | 351-585 |
| U17162 | BAG-1; bcl-2 binding protein with anti-cell death activity | C1e | 17-334 |
| U15784 | Shc transforming adaptor protein; Src homology 2 (SH2) protein, SHBrelated | A5t | 1220-1451 |
| U18310 | MAPKK4; MAP kinase kinase 4; Jnk activating kinase 1; (JNKK1; SEK1; MKK4) | B6c | 1380-1749 |
| U19118 | Transcription factor LRG - 21 | D6n | 618-966 |
| U19119 | Interferon inducible protein 1 | D4k | 1342-1636 |
| U19463 | A20 zinc finger protein; apoptosis inhibitor | C2e | 1952-2293 |
| U19596 | p18ink4; cdk4 and cdk6 inhibitor | A7c | 16-284 |
| U19799 | 1-kB (l-kappa B) beta | B3n | 419-778 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| U24160 | Dvi2; dishevelled-2 tissue polarity protein | B7i | 1205-1578 |
| U20532 | Nuclear factor related to P45 NF-E2 | D5h | 1429.1759 |
| U21011 | MSH2 DNA mismatch repair protein; MutS homologue 2 | C7a | 2150-2490 |
| U20238 | Gaplli; GTPase-activating protein | B7i | 328-644 |
| U25685 | Syk tyrosine-protein kinase (activated p21cdc42Hs kinase (ack)) | B5d | 1235-1524 |
| U27177 | p107; RBL1; Retinoblastoma gene product-related protein p107 (cell cycle regulator) | A1j | 1973-2365 |
| U28724 | PMS2 DNA mismatch repair protein; yeast PMS1 homolog 2 | C7d | 749-1013 |
| U29173 | Limphotoxin receptor (TNFR family) | E2g | 1415-1668 |
| U31625 | BRCA1; BreasVovarian cancer susceptibility locus 1 product | A1b | 5126-5430 |
| U33626 | Pml; Murine homologue of the leukemia-associated PML gene | B4b | 1667-2064 |
| U34960 | Transducin beta-2 subunit | B7e | 515.834 |
| U36277 | 1-kB (1-kappa B) alpha chain | B3m | 541-823 |
| U37522 | TRAIL; TNF-related apoptosis inducing ligand; Apo-2 ligand | C5c | 981-1288 |
| U36799 | p130; Retinoblastoma gene product-related protein Rb2/p130 (cell cycle regulator) | A1k | 970-1321 |
| U36340 | CACCC Box- binding protein BKLF | D1j | 826-1065 |
| U39643 | FAF1; Fas-associated protein factor, apoptosis activator | C3e | 423.681 |
| U41671 | Zinc finger transcription factor RU49 | D7m | 1229-1591 |
| U42190 | GTBP; G/T-mismatch binding protein; MSH6 | C6g | 1477-1769 |
| U43144 | PLC beta; phospholipase C beta 3 | B6I | 1933-2271 |
| U43205 | Frizzled-3; Drosophila tissue polarity gene frizzled homologue 3; dishevelled receptor | B2m | 2037-2285 |
| U43187 | MAPKK3; MAP kinase kinase 3 (dual specificity) (MKK3, MEK3) | B6b | 1436-1742 |
| Ü43525 | Myeloblastin; trypsin-chymotrypsin related serine protease | A71 | 503-807 |
| U47104 | Zinc finger Kruppel type Zip 92 | D71 | 578-896 |
| U44088 | TDAG51; couples TCR signaling to Fas (CD95) expression | C5a | 729-1042 |
| U43788 | POU domain, class 2, associated factor 1 | D6c | 610-884 |
| U48853 | Cas; Crk-associated substrate; focal adhesion kinase substrate | B41 | 1982-2216 |
| U49112 | ALG-2; calcium binding protein required for programmed cell death | C2i | 527-861 |
| U49739 | Unconventional myosin VI | F6e | 3784-4021 |
| U51037 | Transcription factor CTCF (11 zinc fingers) | D61 | 1625-1911 |
| U53925 | Transcription factor C 1 | D6k | 3895-4227 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| U58992 | Madr1; mSmad1; Mothers against dpp protein (Mad) murine homologue; TGF-beta signaling protein-1 (bsp-1); candidate tumor suppressor gene | A1g | 238-476 |
| U59746 | Bcl-W apoptosis regulator; $\mathrm{Bcl}-2$ family member | C1i | 153-368 |
| U60530 | Mad related protein 2 (MADR2) | F3h | 584-820 |
| U62638 | Cyclin C (G1-specific) | A6e | 714-986 |
| U63386 | Mph-1 nuclear transcriptional repressor for hox genes | D5a | 1621-1884 |
| U66887 | Rad50; DNA repair protein | C71 | 1383-1707 |
| U70324 | Fyn proto-oncogene; Src family member | B5a | 584-882 |
| X01023 | c-myc proto-oncogene protein | A21 | 379-667 |
| V00727 | c-Fos proto-oncogene; transcription factor AP-1 component. fos cellular oncogene | A2h | 482.734 |
| $\times 06086$ | Cathepsin L | F6j | 267.588 |
| X04648 | Glutamate receptor channel subunit gamma | E6n | 41-408 |
| X12616 | c-Fes proto-oncogene | A41 | 2342-2598 |
| X12822 | Cytotoxic cell protease 2 (B10) | F61 | 439-686 |
| X07439 | Homeo Box protein 3.1 (Hox-3.1) | D4d | 449-722 |
| X13721 | Homeo Box protein 2.4 (Hox-2.4) | D4b | 1949-2284 |
| X14897 | Fos-B; c-fos-related protein fos B | A3c | 920-1278 |
| X16490 | Plasminogen activator inhibitor-2 | F7i | 674.978 |
| X51983 | c-ErbA oncogene; thyroid hormone receptor. | A2g | 400-675 |
| X 53337 | Cathepsin D | F6h | 587-894 |
| $\times 51438$ | Vimentin | F6d | 868-1096 |
| $\times 53476$ | HMG-14 non histone chromosomal protein | D3m | 643-1017 |
| X53798 | Macrophage inflamatory protein 2 alpha (MIP 2 alpha) | F3g | 14-352 |
| X56906 | Bone morphogenetic protein 7 (BMP-7) (osteogenic protein 1) | F1e | 670-971 |
| X56959 | Transcription factor SP1P (POUdomain transcription factor) | D71 | 866-1128 |
| X59252 | Homeo Box protein 8 (Hox-8) | D4g | 826-1132 |
| X59927 | Fibroblast growth factor receptor 4 | E2b | 2446-2820 |
| X57277 | Rac1 murine homologue | B7c | 425-651 |
| X60831 | Transcription factor UBF | D7h | 689-993 |
| $\times 61435$ | Kinesin heavy chain | F5n | 1898-2182 |
| $\times 61800$ | CCAAT- Binding transcription factor (C/ EBP) | D1k | 904-1150 |
| X62622 | TIMP-2 tissue inhibitor of metalloproteinases-2 | F7m | 1236-1468 |
| $\times 63190$ | Ets-related protein PEA 3 | D3a | 1702-2040 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| X64361 | Vav; GDP-GTP exchange factor; proto-oncogene | B71 | 1083-1351 |
| X63963 | PAX-6 (paired box protein) | D6b | 1081-1325 |
| X66032 | Cyclin B2 (G2/M-specific) | A6d | 874-1236 |
| $\times 67083$ | Chop10; murine homologue of Gadd153 (growth arrest and DNA-damageinducible protein) | C3a | 17-332 |
| X67914 | PD-1 possible cell death inducer; Ig gene superfamily member | C4i | 1481-1734 |
| X69619 | Inhibin beta A subunit (TGF beta family) | F2h | 1064-1304 |
| $\times 70842$ | Vegir2; KDR/flk1 vascular endothelial growth factor tyrosine kinase receptor | B3j | 1394-1721 |
| X70296 | Protease nexin 1 (PN-1) | F7d | 746-985 |
| X71327 | MRE-binding transcription factor | D5b | 552-916 |
| X72711 | Activator -1 140 KD subunit (replication factor C 140KD) | C5e | 4137-4375 |
| X72310 | DP. 1 (DRTF-polipeptide 1) cell cycle regulatory transcription factor | D2g | 925-1305 |
| X72230 | 5-Hydroxytryptamine (serotonin) receptor 1c | E49 | 982-1314 |
| $\times 72795$ | Gelatinase B | F6n | 599-954 |
| X74351 | XPAC; xeroderma pigmentosum group $A$ correcting protein | C7m | 447-669 |
| X75427 | Integrin alpha 2 (CD49b) | E7a | 1595-1976 |
| X77113 | Growh/ differentiation factor 2 (GDF-2) | F2c | 939-1329 |
| X81582 | Insulin-like growth factor binding protein-4 (IGFBP-4) | F21 | 781-1140 |
| X81579 | Insulin-like growth factor binding protein-1 (IGFBP-1) | F2i | 27-256 |
| $\times 81580$ | IGFBP-2; insulin-like growth factor binding protein 2; autocrine and/or paracrine growth promoter | A5m | 449-817 |
| X81583 | Insulin-like growth factor binding protein-5 (IGFBP-5) | F2m | 461-824 |
| X81584 | Insulin-like growth factor binding protein -6 (IGFBP 6) | F2i | 701-1039 |
| X82327 | A-myb proto-oncogene; myb-related protein A | A2e | 1017-1334 |
| X83536 | Membrane type matrix matallóproteinase | F7c | 877.1101 |
| X87257 | Elk-1 ets-related proto-oncogene | A3a | 1498-1680 |
| X86925 | E2F-5 transcription factor | D2h | 426-728 |
| X90829 | Lbx 1 transcription factor | D4n | 1000-1306 |
| X91144 | P-selectin (glycoprotein ligand-1) | E51 | 1095-1323 |
| X91753 | Transcription factor SEF2 | D7e | 755-1054 |
| Z11974 | Macrophage mannose receptor | E2h | 807-1197 |
| X95403 | Rab-2 ras-related protein | B7b | 232-505 |
| X98055 | Gluthathione S-transferase (theta type1); phase II conjugation enzyme | C2c | 14-298 |
| X99063 | Zyxin; LIM domain protein; alpha-actinin binding protein | B7n | 1437-1812 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| Y00671 | Met protooncogene | A4d | 3646-3933 |
| Y00864 | c-Kit proto-oncogene (maststem cell growth factor receptor tyrosine kinase) | A4c | 2867-3181 |
| Y07960 | Transcription factor BARX1 (homeodian transcription factor) | D6j | 723.973 |
| X95346 | PLC gamma; phospholipase C gamma | B6m | 180-516 |
| Z12604 | Stromelysin-3; matrix metalloproteinase-11 (MMP-11) | C4n | 1463-1806 |
| 214224 | 5-Hydroxytryptamine (serotonin) receptor 1e beta | E4h | 530-774 |
| 215119 | 5-Hydroxytryptamine (serotonin) receptor 2c | E4i | 588-940 |
| 219521 | Low density lipoprotein receptor | E4d | 1047-1324 |
| Z23107 | 5.Hydroxytryptamine (serotonin) receptor 7 | E4k | 460-817 |
| 222649 | c-Mpl; thrombopoietin receptor; hematopoietic growth factor receptor superfamily member | A5k | 1561-1772 |
| Z21848 | DNA-polymerase delta catalytic subunit | C6b | 1256-1600 |
| 229532 | Follistatin | F11 | 764-1053 |
| 247766 | Cyclin F (S/G2/M-specific) | A6j | 2431-2708 |
| 236885 | Ets-related protein Sap 1A | D3c | 1267-1521 |
| Z32815 | Net; ets related transcription factor; activated by Ras | A3i | 1211-1595 |
| Z48538 | Stat5a; mammary gland factor | B4f | 2269-2628 |
| 249086 | Hek2 murine homologue; Mdk5 mouse developmental kinase; Eph -related tyrosine-protein kinase receptor | B2n | 1702-1930 |
| D26177 | D-Factor/LIF receptor | E11 | 2376-2775 |
| M13806 | Cytoskeletal epidermal keratin (14 human) | F5h | 108-469 |
| M21019 | R-ras protein, closely related to ras proto-oncogenes | B7d | 215-555 |
| M22959 | Prolactin receptor PRLR2 | E4c | 1 1-328 |
| M30903 | Blk; B lymphocyte kinase; Src family member | C2j | 1307-1672 |
| M35590 | Macrophage inflamatory protein 1 beta (Act 2) | F3i | 119-445 |
| M75716 | Alpha-1 protease inhibitor 2 | F7g | 625-969 |
| M92378 | GABA-A transporter 1 | E5e | 1131-1416 |
| M97017 | Bone morphogenetic protein 8a (BMP-8a) (TGF-beta family) | F1f | 788-1139 |
| M97200 | Erythroid kruppel-like transcription factor | D2n | 783.1171 |
| M98339 | GATA binding transcription factor (GATA-4) | D3e | 81.379 |
| M98547 | Growth factor receptor | E2f | 1701-2014 |
| S72408 | Crk adaptor protein | B4m | 750-1027 |
| U09419 | Retinoid X receptor interacting protein (RIP 15) | D6g | 1388-1682 |
| U14752 | Cek 7 receptor protein tyrosine kinase ligand | F1h | 504-837 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| U29678 | C-C CKR-1; CCR-1; C-C chemokine receptor type 1, macrophage inflammatory protein-1 alpha receptor; MIP-1alpha-R; RANTES-R | B2i | 168-495 |
| X13358 | Glucocorticoid receptor form A | E3m | 1527-1816 |
| $\times 83106$ | Mothers against DPP protein (mad homolog Smad 1, transforming growth factor beta signaling protein) | F3j | 464-728 |
| Y00487 | Hck tyrosine-protein kinase | B5b | 1308-1563 |
| AB000777 | Photolyase/blue-light receptor homologue | C7c | 1418-1737 |
| 049482 | Osp94 osmotic stress protein; APG-1; hsp70-related | B1f | 1026-1266 |
| D78645 | Glucose regulated protein, 78 kD ; Grp78 | B1m | $167-411$ |
| D87747 | LCR-1; CXCR-4; CXC (SDF-1) chemokine receptor 4; HIV coreceptor (fusin): G protein-coupled receptor LCR1 homologue; | B3d | 584-867 |
| M23384 | Glucose transporter-1, erythrocyte; Glut1 | B2e | 325-653 |
| M80456 | Int-3 proto-oncogene; NOTCH family member; NOTCH4 | A5h | 1846-2145 |
| M94335 | c-Akt proto-oncogene; Rac-alpha; proteine kinase B (PKB) | C2k | 604-899 |
| Y13231 | Bak apoptosis regulator; Bcl-2 family member | C11 | 1509-1786 |
| U57324 | PS-2; homologue of the Alzheimer's disease gene | C4h | 437.783 |
| U65594 | BRCA2; Breast cancer susceptibility locus 2 product | A1c | 649-922 |
| U66058 | DNA ligase III | C5k | 2980-3205 |
| U67321 | Caspase-7: Lice2; ICE-LAP3 cysteine protease | C1c | 1040-1280 |
| U75506 | BID; apoptic death agonist | C1k | 452.777 |
| U92456 | WBP6; pSK-SRPK1; WW domain binding protein 6 serine kinase for SR splicing factors | B7m | 482.774 |
| U95826 | Cyclin G2 (G2/M-specific) | A6I | 408-688 |
| $\times 99018$ | Ung1; uracil-DNA glycosylase | C71 | 444-729 |
| Y14019 | Rab-3b ras-related protein | F6c | 232-562 |
| U28423 | Inhibitor of the RNA-activated protein kinase, $58-\mathrm{kDa}$ | B5i | 180.487 |
| U34259 | Golgi 4-transmembrane spanning transporter; MTP | B2d | 742-1060 |
| U34920 | ATP-binding casette 8; ABC8; homolog of Drosophila white | B2b | 1011-1319 |
| U37720 | CDC42 GTP-binding protein; G25K | F5g | 1675-1982 |
| U41751 | Etoposide induced p53 responsive (EI24) mRNA | B11 | 1041-1296 |
| U51866 | Casein kinase II (alpha subunit) | A3n | 1237-1517 |
| U52945 | TSG101 tumor susceptibility protein | A1n | 446-713 |
| U54705 | Tumor suppressor maspin | A2a | 251.507 |
| U97076 | FLIP-L; apoptosis inhibitor; FLICE-like inhibitory protein | C3h | 1476-1811 |
| Х63615 | CamK II; Ca2+/calmodulin-dependent protein kinase II (beta subunit) | F5f | 1951-2219 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| 249085 | Htk; Mdk2 mouse developmental kinase; Eph -related tyrosine-protein kinase receptor | B3a | 2032-2365 |
| D49921 | Glial cell line-derived neurotrophic factor | F1n | 236-539 |
| L06039 | CD31 (Platelet endothelial cell adhesion molecule 1) | E6d | 1172-1494 |
| L16928 | CD22 antigen | E6i | 2314-2645 |
| L39770 | Gbx 2 | D3g | 1122-1395 |
| M12302 | Cytotoxic T lymphocyte-specific serine protease CCP I gene (CTLA-1) | F6m | 585-830 |
| M14222 | Cathepsin B | F6g | 382-729 |
| M33324 | Growth hormone receptor | E3n | 1942-2240 |
| M34563 | CD28 (receptor for B71) | E6b | 544-774 |
| M38651 | Estrogen receptor | E31 | 742-1013 |
| S71251 | Monotype chemoattractant protein 3 | E1k | 201-491 |
| U03856 | CD45 associated protein (CD 45-ap, LSM-1) | E6f | 620-898 |
| U11688 | Orphan receptor | E1b | 1686-1943 |
| U17985 | Cannabinoid receptor 1 (brain) | E4n | 1091-1437 |
| U43512 | Dystroglycan 1 | E6m | 2267-2505 |
| U46923 | G-protein coupled receptor | E5c | 350-671 |
| X02389 | Urokinase type plasminogen activator | F7f | 1301-1538 |
| X05719 | CTLA-4 (immunoglobin superfamily member) | E6k | 246-519 |
| X56182 | Myogenic factor 5 | D5d | 232-528 |
| X62700 | UPAR1; urokinase plasminogen activator surface receptor (CD87) | B3i | 482.756 |
| X69832 | Serine protease inhibitor 2.4 | F7k | 621-927 |
| X70298 | SRY-box containing gene 4 | D7b | 34-311 |
| L25602 | Bone morphogenetic protein 2 (BMP-2) (TGF-beta family) | F1c | 8372-8724 |
| M10021 [K02 | [K02588] P-1-450; dioxin-inducible cytochrome P450 | B2a | 3729-4014 |
| M16506 | Bcl-2; B cell lymphoma protein 2, apoptosis inhibitor | C1h | 2125-2367 |
| M34510 | CD14 antigen | E6h | 667.931 |
| M81832 | Somatostatin receptor 2 | E3b | 47-310 |
| U19880 | Dopamine receptor 4 | E5b | 907-1191 |
| U21681 | Cannabinoid receptor 2 (macrophage, CB2) | E5a | 910-1262 |
| U58533 | Enf (Ets-related transcription factor) | D2m | 1286-1613 |
| 211597 | 5-Hydroxytryptamine (serotonin) receptor 1b | E4if | 1043-1355 |
| D78382 | Tob antiproliferative factor; interacts with p185erbB2 | A7n | 540-876 |
| J03752 | Glutathione S-transierase (microsomal) | C2a | 185-428 |
| L20331 | Adenosine A3 receptor | C2h | 182-382 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| U05341 | p55cdc; cell division control protein 20 | C4e | 1061-1348 |
| U12273 | AP endonuclease; apurinic/apyrimidinic endonuclease (Apex) | C5f | 1894-2150 |
| $\times 67735$ | Mas proto-oncogene (G-protein coupled receptor) | A51 | 566-808 |
| D26046 | AT motif-binding factor ATBF1 | D1d | 9807-10112 |
| D49474 | HMG-box transcription factor from testis (MusSox17) | D31 | 427-662 |
| L03547 | Ikaros DNA binding protein | D4i | 627-890 |
| L12147 | Early B cell factor (EBF) | D2a | 750-1026 |
| L12703 | Engrailed protein (En-1) homolog | D2b | 1323-1554 |
| L12705 | Engrailed protein (En-2) homolog | D2c | 1626-1895 |
| L21027 | Transcription factor A10 | B4i | 499-806 |
| L26507 | Myocyte nuclear factor (MNF) | D5c | 1203-1456 |
| L36435 | Basic domain/leucine zipper transcription factor | D1e | 872-1073 |
| M37163 | Caudal type Homeobox 1 (Cdx1) | D11 | 1040-1301 |
| M58566 | Butyrate response factor 1 | D1i | 768-1054 |
| S53744 | Brain specific transcription factor NURR-1 | D19 | 1548-1754 |
| S68377 | Brn-3.2 POU transcription factor | D1h | 877-1237 |
| 574520 | Caudal type Homeobox 2 (Cdx2) | D1m | 1085-1367 |
| U01036 | Erythroid transcription factor NF-E2 | D2d | 1-241 |
| U20344 | Gut-specific Kruppel-like factor GKLF | D3i | 1558-1789 |
| U25096 | Kruppel-like factor LKLF | D4m | 898-1193 |
| U29086 | Neuronal helix-loop-helix protein NEX-1 | D5e | 572-907 |
| U36760 | Brain factor 1 (Hfhbf1) | D1f | 1080-1318 |
| U41626 | Split hand/foot gene | D5m | 92-303 |
| U42554 | Sim transcription factor | D1n | 2828-3066 |
| U59876 | Glial cells missing gene homolog (mGCM1) | D3h | 727-1080 |
| U62522 | Sp4 zinc finger transcription factor | D4j | 1704-1929 |
| X61754 | Heat shock transcription factor 2 (HSF 2) | D3j | 1445-1640 |
| $\times 83974$ | RNA polymerase I termination factor TTF-1 | A2j | 3222-3433 |
| L35949 | Hepatocyte nuclear factor 3/forkhead homolog 8 (HFH-8) | D3k | 913-1232 |
| X94125 | SRY-box containing gene 3 (Sox3) | D5n | 212-443 |
| D13759 | Cot proto-oncogene | A3m | 696-956 |
| D49429 | HR21spA; protein involved in DNA double-strand break repair; PW29; calcium-binding protein | C6h | 103-434 |
| D64107 | MmLim15; RecA-like gene; DMC1 homologue; meiosis-specific homologous recombination protein | C6I | 581-781 |


| GenBank \# | Gene Name | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| J 05186 | ERp72 endoplasmic reticulum stress protein; protein disulfide isomeraserelated protein | B1k | 1160-1470 |
| S50213 | HMG1-related VDJ recombination signal binding protein | B1h | 2263-2531 |
| S65038 | Gli oncogene; zinc finger transcription factor | A3e | 104-505 |
| U05245 | Tiam-1 invasion inducing protein; GDP-GTP exchanger-related | A5n | 4329-4628 |
| U16805 | Sik; Src-related intestinal kinase | C4k | 1246-1623 |
| U28495 | Lic proto-oncogene | A5d | 853-1150 |
| U40930 | Oxidative stress-induced protein mRNA | B1n | 1248-1561 |
| U43900 | STAM; signal transducing adaptor molecule | C4m | 576-811 |
| U46854 | ShcC adaptor; Shc-related; brain-specific | C7i | 246-601 |
| U58987 | MmMre11a putative endo/exonuclease | B1i | 866-1204 |
| X53068 | PCNA; proliferating cell nuclear antigen; processivity factor | C7b | 53-320 |
| X81464 | Translin; recombination hotspot binding protein | C7j | 205-431 |
| X96618 | PA6 stromal protein; RAG1 gene activator | C6a | 442-749 |
| U18342 | Sky proto-oncogene (Tyro3; Rse; Dtk) | A4h | 1927-2286 |
| 250013 | H-ras proto-oncogene; transforming G-protein | A5c | 1307-1544 |
| L47239 | ERBB-2 receptor (c-neu, HER2 protein tyrosine kinase) | E1m | 16-42 |
| L47240 | ERBB-3 receptor | E1n | 4-243 |
| U22516 | Placental ribonuclease inhibitor (Angiogenin) | F4a | 512-766 |
| L00923 | myosin 1 | G13 | 2578-2921 |
| U459777 | Ca2+ binding protein, Cab45 | G20 | 597-1082 |
| M10624 | murine ornithine decarboxylase | G14 | 865-1252 |
| $\times 51703$ | ubiquitin | G5 | 123-547 |
| J00423 | Hypoxantine-guanine phosphoribosyltransferase | G7 | 301-751 |
| D78647 | phospholipase A2 | G6 | 446-813 |
| L31609 | ribosomal protein S 29 | G21 | 5-244 |
| M325999 | glyceraldehyde-3-phosphate dehydrogenase | G12 | 765-1016 |
| M12481 | beta-actin | G19 | 25-564 |

## Cancer Array

In the cancer arrays of the subject invention, the polynucleotide probe compositions on the array correspond to those genes which are associated, e.g. play a role in, cellular proliferative diseases, particularly cancer, where human genes are of particular interest in many embodiments. Types of genes that are typically represented on a cancer array of the subject invention include: oncogenes, tumor suppressors, cell cycle regulators, genome plasticity genes, apoptosis genes, cell differentiation genes, regulators of tumor host interaction and metastasis, such as extracellular matrix proteins, cell adhesion receptors, molecules that control cell invasion and motility, and genes associated with angiogenesis.

In certain embodiments, of particular interest is an array having the following types of genes represented on its surface: cell cycle/growth regulators; apoptosis; growth factors/cytokines; oncogenes/tumor suppressors; cell adhesion, motility and invasion; invasion regulators; GTP ases and their regulators; cadherins; intermediate filament markers; receptors; cell fate/development regulators; DNA damage/response/repair/recombination; and angiogenesis regulators. In a specific cancer array of interest, the spots are as listed in Table 3.

The cancer array finds use in a variety of applications, including: monitoring cellular responses to therapeutic compounds; comparing expression profiles of tumors at different developmental stages; developing diagnostic tools for distinguishing closely related tumors; and the like.

In the following Table 3, as well as preceding Tables 1 and 2, the "position" coordinate refers to the actual nucleotide residues of the listed gene that are represented on the array.

|  | Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: | :---: |
|  | - QUADRANT A |  |  |  |
|  | CELL DIVISION CONTROL PROTEIN 2 HOMOLOG (EC 2.7.1.-) (P34 PROTEIN KINASE) (CYCLIN-DEPENDENT KINASE 1) (CDK1) | X05360 | A1a | 655-886 |
|  | CELL DIVISION PROTEIN KINASE 2 (EC 2.7.1.-) (P33 PROTEIN KINASE) | M68520 | A1b | 1774-2180 |
|  | CELL DIVISION PROTEIN KINASE 3 (EC 2.7.1.-). | X66357 | A1c | 216.882 |
|  | CELL DIVISION PROTEIN KINASE 4 (EC 2.7.1.-) (PSK-J3) | M14505 | A1d | 372.693 |
|  | CELL DIVISION PROTEIN KINASE 5 (EC 2.7.1.-) (TAU PROTEIN KINASE II CATALYTIC SUBUNIT) (TPKII CATALYTIC SUBUNIT) (KINASE PSSALRE). | X66364 | A1e | 468-767 |
|  | CELL DIVISION PROTEIN KINASE 6 (EC 2.7.1.-) (KINASE PLSTIRE) | $\times 66365$ | A1f | 315-663 |
|  | CELL DIVISION PROTEIN KINASE 7 (EC 2.7.1.-) (CDK-ACTIVATING KINASE) (CAK) (39 KD PROTEIN KINASE) (P39 MO15) (STK1) (CAK1). | L20320 | A1g | 89-305 |
|  | CYCLIN-DEPENDENT KINASE 5 ACTIVATOR ISOFORM P39I PRECURSOR (CDK5 ACTIVATOR) (P39I). | U34051 | A1h | 763-1-62 |
| $\cdots$ | CYCLIN-DEPENDENT KINASE 5 ACTIVATOR PRECURSOR (CDK5 ACTIVATOR) (TAU PROTEIN KINASE II 23 KD SUBUNIT) (TPKII REGULATORY SUBUNIT) (P23) (P25) (P35). | X80343 | A1i | 551.941 |
| $\underset{\sim}{\sim}$ | cdc25A; M-PHASE INDUCER PHOSPHATASE 1 (EC 3.1.3.48) | M81933 | A1i | 1632-1978 |
| 1 | cdc25B; M-PHASE INDUCER PHOSPHATASE 2 (EC 3.1.3.48). (CDC25Hu2) | M81934; [S78187] | A1k | 2286-2602 |
|  | cdc25C; M-PHASE INDUCER PHOSPHATASE 3 (EC 3.1.3.48). | M34065 | A11 | $331-623$ |
|  | CLK-1 | L29222 | A1m | 144.459 |
|  | CLK-2 | L29216 | A1n | 1106-1356 |
|  | CLK-3 | L29220 | A2a | 551-1002 |
|  | SERINETHREONINE-PROTEIN KINASE KKIALRE | X66358 | A2b | 276.461 |
|  | SERINETHREONINE-PROTEIN KINASE PCTAIRE-1 | X66363 | A2C | 1114-1434 |
|  | SERINETHREONINE-PROTEIN KINASE PCTAIRE-2 | $\times 66360$ | A2d | 954-1250 |
|  | SERINETHREONINE PROTEIN KINASE PCTAIRE-3 | X66362 | A2e | 549-911 |
|  | SERINETHREONINE PROTEIN KINASE PITALRE | L25676 | A21 | $367-635$ |
|  | CDC2-RELATED PROTEIN KINASE CHED | M80629 | A2g | 1388-1548 |
|  | CDC2-RELATED KINASE PISSLRE | L33264 | A2h | 454-755 |
|  | CYCLIN A | X51688 | A2i | 876-1218 |
|  | CYCLIN 81 G2/MITOTIC-SPECIFIC | M25753 | A2j | 979-1311 |
|  | CYCLIN C G1/S-SPECIFIC | M74091 | A2k | 6670-7326 |
|  | CYCLIN D1 (CYCLIN PRAD1) (BCL-1 ONCOGENE) | X59798; [M64349] | A21 | 3427-3784 |
|  | CYCLIN D2 | D13639 (M90813) | A2m | 3932.4284 |
|  | CYCLIN D3 | M92287 | A2n | 537-894 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| CYCLIN E | M73812 | A3a | 1295-1658 |
| CYCLIN G1 | U47413 [L49504] | A3b | 755-1035 |
| CYCLIN G2 | U47414 [L49506] | A3C | 989-1254 |
| CYCLINH | U11791 [U12685] | A3d | 717-1026 |
| CYCLIN-DEPENDENT KINASE INHIBITOR 1 (MELANOMA DIFFERENTIATION ASSOCIATED PROTEIN 6) (MDA-6) (P21) (CDKINTERACTING PROTEIN 1) (CIP1) (WAF1) (CDKN1A) (CDKN1) (SDI1) (PIC1) (CAP20) | U09579; [L25610] | A3e | 1745-2063 |
| CYCLIN-DEPENDENT KINASE INHIBITOR 1C (CYCLIN-DEPENDENT KINASE INHIBITOR P57) (P57KIP2) | U22398 | A3f | 1048-1316 |
| CYCLIN-DEPENDENT KINASE 4 INHIBITOR A (CDK4I) (P16-INK4) (P16INK4A) (MULTIPLE TUMOR SUPPRESSOR 1) (MTS1). (CDKN2A) | L27211 | A3g | 482-836 |
| CYCLIN-DEPENDENT KINASE 4 INHIBITOR B (P14-INK4B) (P15-INK4B) (MULTIPLE TUMOR SUPPRESSOR 2) (MTS2) (CDKN2B). | U17075; [L36844] | A3h | 116-462 |
| CYCLIN-DEPENDENT KINASE 4 INHIBITOR D (P19-INK4D). | U40343; [U20498] | A3i | 750-952 |
| WEE1-LIKE PROTEIN KINASE (EC 2.7.1.112) (Wee1Hu) | U10564 | A3j | 1259-1502 |
| SERINETHREONINE-PROTEIN KINASE PLK (EC 2.7.1.-) (PLK-1) (STPK13) | U01038 | A3k | 1330-3233 |
| PHOSPHOLIPASE D1 | U38545 | A3I | 2862-3961 |
| NEDD5 PROTEIN HOMOLOG. | D63878 | A3m | 381.675 |
| CDC10 PROTEIN HOMOLOG | 572008 | A3n | 66-379 |
| CDC27HS PROTEIN | U00001 | A4a | 870-3474 |
| UBIQUITIN-CONJUGATING ENZYME E2-CDC34 | L22005 | A4b | 249-550 |
| CDC16HS. | U18291 | A4c | 45-378 |
| CDC37 HOMOLOG. | U63131 | A4d | 519.1464 |
| CDC6-RELATED PROTEIN | U77949 | A4e | 216-447 |
| EXTRACELLULAR SIGNAL-REGULATED KINASE 1 (EC 2.7.1.-) (ERK1) (INSULIN- STIMULATED MAP2 KINASE) (MAP KINASE 1) (MAPK 1) (P44 ERK1) (ERT2) (P44-MAPK) (MICROTUBULE-ASSOCIATED PROTEIN-2 KINASE). | X60188 | A4f | 754-1094 |
| EXTRACELLULAR SIGNAL-REGULATED KINASE 3 (EC 2.7.1.-) (ERK3) (MAP KINASE ISOFORM P97) (P97-MAPK). | X80692 | A4g | 806-1267 |
| EXTRACELLULAR SIGNAL-REGULATED KINASE 4 (EC 2.7.1.-) (ERK4) (MAP KINASE ISOFORM P63) (P63-MAPK). | X59727 | A4h | 2678-2994 |
| EXTRACELLULAR SIGNAL-REGULATED KINASE 5 (EC 2.7.1.-) (ERK5) (ERK4) (BMK1 KINASE) | U25278 | A4i | 1010-1267 |
| EXTRACELLULAR SIGNAL-REGULATED KINASE 6 (EC 2.7.1.-) (ERK6) (ERK5) | X79483 | A4j | 530-831 |


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| :---: | :---: | :---: | :---: |
| MITOGEN-ACTIVATED PROTEIN KINASE P38 (EC 2.7.1.-) (MAP KINASE P38) (CYTOKINE SUPPRESSIVE ANTI-INFLAMMATORY DRUG BINDING PROTEIN) (CSAID BINDING PROTEIN) (CSBP) (MAXINTERACTING PROTEIN 2) (MAP KINASE MXI2). | L35253; [L35263] | A4k | 925-1204 |
| STRESS-ACTIVATED PROTEIN KINASE JNK1 (EC 2.7.1.-) (C-JUN NTERMINAL KINASE 1) (JNK-46) | L26318 | A41 | 952-1263 |
| STRESS-ACTIVATED PROTEIN KINASE JNK2 (EC 2.7.1.-) (C-JUN NTERMINAL KINASE 2) (JNK-55). | L31951 | A4m | 638-1000 |
| STRESS-ACTIVATED PROTEIN KINASE JNK3 (EC 2.7.1.-) (C-JUN NTERMINAL KINASE 3) (JNK3) (MAP KINASE P49 3F12). | U34819; [U07620] | A4n | 1018-1413 |
| DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 5 (EC 2.7.1.-) (MAP KINASE KINASE 5) (MAPKK 5) (MAPKJERK KINASE 5). | U25265 | A5a | 629-847 |
| DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 1 (EC 2.7.1.-) (MAP KINASE KINASE 1) (MAPKK 1) (ERK ACTIVATOR KINASE 1) (MAPKERK KINASE 1) (MEK1). | L05624 | A5b | 842-1217 |
| DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 6 (EC 2.7.1.-) (MAP KINASE KINASE 6) (MAPKK 6) (MAPKJERK KINASE 6) (SAPKK3) | U39657 | A5c | 1060-1389 |
| MEK KINASE 3 | U78876 | A5d | 1195-1453 |
| PCNA (CYCLIN) | M15796; (J04718) | A5e | 157.436 |
| PIN1 | U49070 | A5 | 624-1075 |
| RBP1(RETINOBLASTOMA-BINDING PROTEIN) | S57153; S57160 | A5g | 2676-2889 |
| E2F. 1 pRB-binding protein | M96577 | A5h | 899-1595 |
| E2F-3 | Y10479 | A5i | 698-897 |
| E2F-5 | U15642 | A5j | 645-922 |
| E2F-related transcription factor (DP-1) | L23959 | A5k | 935-1186 |
| DP2 (Humdp2) , dimerization partner of E2F | U18422 | A5I | 1603-1838 |
| RBQ-3 | $\times 85134$ | A5m | 359.603 |
| GROWTH-ARREST-SPECIFIC PROTEIN 1 (GAS-1). | L13698 | A5n | 1550-1701 |
| growth inhibitor p33ING1 (ING1) | AF001954 | A6a | 722.983 |
| Abl interactor 2 (Abi-2) + Abl binding protein 3 (AbBP3) [ArgBPIB] | U23435; U31089 | A6b | 1049-1203 |
| GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2 ADAPTOR PROTEIN) (ASH PROTEIN). | L29511; [M96995] | A6c | 355-573 |
| GRB-IR / GRB10 | U69276 | A6d | 358-1155 |
| RAF ONCOGENE | X03484 | A6e | 1704-1989 |
| ral, b- | M95712 | A6i | 866-1144 |
| jun B TRANSACTIVATOR | M29039 | A6g | 1197-1442 |
| N -myc | M13228 | A6h | 761-1188 |


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| :---: | :---: | :---: | :---: |
| C-myc binding protein | D89667 | A6i | 218-490 |
| INTERMEDIATE FILAMENT MARKERS |  |  |  |
| KERATIN, TYPE I CYTOSKELETAL 9 (CYTOKERATIN 9) (K9) (CK 9). | Z29074; [S69510] | A6j | 652-1781 |
| KERATIN, TYPE I CYTOSKELETAL 10 (CYTOKERATIN 10) (K10) (CK 10) | M19156 | A6k | 295-497 |
| KERATIN, TYPE I CYTOSKELETAL 12 (CYTOKERATIN 12) (K12) | D78367 | A6I | 455-624 |
| KERATIN, TYPE I CYTOSKELETAL 13 (CYTOKERATIN 13) (K13) (CK 13) +KERATIN, TYPE I CYTOSKELETAL 15 (CYTOKERATIN 15) (K15) (CK 15) +KERATIN, TYPE I CYTOSKELETAL 17 (CYTOKERATIN 17) (K17) (CK 17) (39.1) | X52426; X07696; X62571 | A6m | 383-1001 |
| KERATIN, TYPE I CYTOSKELETAL 14 (CYTOKERATIN 14)(K14) (CK 14) | J00124 | A6n | 339-839 |
| KERATIN, TYPE I CYTOSKELETAL 16 (CYTOKERATIN 16)(K16) (CK 16):pseudo-keratin K16 type I | M21772; M20336 | A7a | 32-522 |
| KERATIN, TYPE I CYTOSKELETAL 18 (CYTOKERATIN 18) (K18) (CK 18) | M26326 | A7b | 706-971 |
| KERATIN, TYPE I CYTOSKELETAL 19 (CYTOKERATIN 19) (K19) (CK 19). | Y00503 | A7c | 726-1124 |
| KERATIN, TYPE II CYTOSKELETAL 1 (CYTOKERATIN 1) (K1) (CK 1) (67 KD CYTOKERATIN) (HAIR ALPHA PROTEIN) | M98776 | A7d | 894-1459 |
| KERATIN, TYPE II CYTOSKELETAL 2 ORAL (CYTOKERATIN 2P) (K2P) (CK 2P) | M99063 | A7e | 2167-2455 |
| KERATIN, TYPE II CYTOSKELETAL 2 EPIDERMAL (CYTOKERATIN 2E) (K2E) (CK 2E) | M99061 [S43646] | A71 | 1091-1450 |
| KERATIN, TYPE II CYTOSKELETAL 4 (CYTOKERATIN 4) (K4) (CK4) | X67683 | A7g | 66.404 |
| KERATIN, TYPE II CYTOSKELETAL 5 (CYTOKERATIN 5) (K5) (CK 5) (58 KD CYTOKERATIN) | M21389 | A7h | 93-682 |
| KERATIN, TYPE II CYTOSKELETAL 6 (CYTOKERATIN 6A) (CK 6A) (K6A KERATIN) +(CYTOKERATIN 6B) (CK 6B) (K6B KERATIN) + (CYTOKERATIN 6C) (CK 6C) (K6C KERATIN) + (CYTOKERATIN 6D) (CK 6D) (K6D KERATIN) + (CYTOKERATIN 6E) (CK 6E) (K6E KERATIN) + (CYTOKERATIN 6F) | $\begin{aligned} & \text { J00269; V01516; L42592; } \\ & \text { L00205; L42601; L42610; } \\ & \text { L42611; L42612 } \end{aligned}$ | A7i | 689-880 |
| KERATIN, TYPE II CYTOSKELETAL 6B (CYTOKERATIN 6B) (CK 6B) (K6B KERATIN) | L42592; L00205 | A7j | 275-414 |
| KERATIN, TYPE II CYTOSKELETAL 7 (CYTOKERATIN 7) (K7) (CK 7) | X03212 | A7k | 1154.1430 |
| KERATIN, TYPE II CYTOSKELETAL 8 (CYTOKERATIN 8) (K8) (CK 8) | M34225 | A71 | 1190-1474 |
| VIMENTIN | X56134 [M14144] | A7m | 460-740 |
| DESMIN | U59167 | A7n | 1063-1364 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| QUADRANT B |  |  |  |
| APOPTOSIS |  |  |  |
| BCL2 | M14745 | B1a | 5078-5382 |
| Bcl2 and p53 binding protein Bbp/53BP2 (BBP/53BP2) | U58334 | B1b | 3129-3376 |
| BAX | L22474 | B1C | 227.478 |
| APOPTOSIS REGULATOR BCL-W | U59747 | B1d | 121.403 |
| INDUCED MYELOID LEUKEMIA CELL DIFFERENTIATION PROTEIN MCL-1 (ORF is at nt. 61-1053; ML) | L08246 | B1e | 697.977 |
| BCL2-RELATED PROTEIN A1 (BFL-1 PROTEIN) (HEMOPOIETICSPECIFIC EARLY RESPONSE PROTEIN) (GRS PROTEIN) | U29680 | B1f | 64-293 |
| BCL-2 INTERACTING KILLER (APOPTOSIS INDUCER NBK) (BP4) (BIP1) (BIK) | X89986; [U34584] | B19 | 935-1200 |
| BCL-2 HOMOLOGOUS ANTAGONISTKILLER (APOPTOSIS REGULATOR BAK) | $\begin{aligned} & \text { U23765; [U16812; } \\ & \text { U16811; } \times 84213 \text { ] } \\ & \hline \end{aligned}$ | B1h | 1371-1661 |
| BAD PROTEIN (BCL-2 BINDING COMPONENT 6). | U66879 | B1i | 408.749 |
| BCL-2 BINDING ATHANOGENE-1 (BAG-1) (GLUCOCORTICOID RECEPTOR-ASSOCIATED PROTEIN RAP46). | S83171; [Z35491] | B1j | 511.830 |
| serine/threonine protein kinase, NIK; binds speciilically to TRAF2 | Y10256 | B1k | 3776-4036 |
| Casper, a FADD- and caspase-related inducer of apoptosis [CASH-alpha+ CASH-beta) (FLAME-1) (FLICE-like inhibitory protein) | AF010127(Y14039; <br> Y14040) | B11 | 363-787 |
| death domain containing protein CRADD, apoptotic adaptor molecule for caspase-2 and FasLTNF receptor-interacting protein RIP | U84388 | B1m | 369-604 |
| TNF receptor-1 associated protein (TRADD) | L41690 | B1n | 1009-1313 |
| cell death protein kinase RIP | U25994; [U50062] | B2a | 848-1123 |
| DAXX, a FAS-binding protein that activates JNK and apoptosis | AF015956 | B2b | 804-1030 |
| Apo-2 ligand (TNF-related apoptosis inducing ligand TRAIL) | U57059 | B2c | 211.616 |
| TRAF-INTERACTING PROTEIN I-TRAF (TRAF family member-associated NF-kB activator TANK) | U59863; [U63830] | B2d | 674-887 |
| TRAF5 | U69108 | B2e | 1318-1694 |
| TRAFE | U78798; [L81153] | B2f | 1689-1961 |
| TRAF-interacting protein (TRIP) | U77845 | B2g | 154-387 |
| tumor necrosis factor type 2 receptor associated protein (TRAP3) | U12597 | B2h | 1207-1566 |
| CD40 RECEPTOR ASSOCIATED FACTOR 1 (CRAF1) (CAP-1), (LMP1 associated protein) | $\begin{aligned} & \text { U21092; [U15637; L38509; } \\ & \text { U19260) } \end{aligned}$ | B2i | 980-1322 |
| INHIBITOR OF APOPTOSIS PROTEIN 1 (HIAP1) (HIAP-1) (C-IAP2) (TNFR2-TRAF SIGNALLING COMPLEX PROTEIN 1) (IAP HOMOLOG C) (IAP1) (MIHC). | U45878; [U37546] | B2j | 1444-1848 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| INHIBITOR OF APOPTOSIS PROTEIN 2 (HIAP2) (HIAP-2) (C-IAP1) (TNFR2-TRAF SIGNALLING COMPLEX PROTEIN 2) (IAP HOMOLOG B) (IAP2) (MIHB). | U45879; [U37547] | B2k | 266-621 |
| X-LINKED INHIBITOR OF APOPTOSIS PROTEIN (X-LINKED IAP) (IAPLIKEPROTEIN) (HILP). | U45880; [U32974] | B21 | 2000-2363 |
| p53-dependent cell growth regulator CGR19 | U66469 | B2m | 28-301 |
| cytotoxic ligand TRAIL receptor | U90875 | B2n | 290-548 |
| (ICE) (INTERLEUKIN-1 BETA CONVERTING ENZYME) (P45) (CASPASE 1) | U13699; [M87507; X65019] | B3a | 5078-5282 |
| (CASPASE-2) (ICH-1L) (ICH-1S) | U13021; [U13022) | B3b | 851-1218 |
| APOPAIN PRECURSOR (EC 3.4.22.-) (CYSTEINE PROTEASE CPP32) (YAMA PROTEIN) (CASPASE-3) (CPP32) (YAMA PROTEIN) (CASPASE- <br> 3) isoform alpha | U13737 | B3c | 2007-2434 |
| ICH-2 PROTEASE PRECURSOR (EC 3.4.22.-) (TX PROTEASE) (ICEREL <br> II) (CASPASE-4) + CASPASE-5 PRECURSOR (EC 3.4.22.-) (ICH-3 PROTEASE) (TY PROTEASE (ICEREL-III). | U28014; U28015 | B3d | 763-11-07 |
| CASPASE-6 PRECURSOR (EC 3.4.22.-) (APOPTOTIC PROTEASE MCH- <br> 2) isoform beta + isolorm alpha | U20537; U20536 | B3e | 387-697 |
| CASPASE-7 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 3) (ICE-LAP3) (APOPTOTIC PROTEASE MCH-3) (CMH-1) (Lice2) | U37448 | B3i | 1042-1413 |
| CASPASE-8 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 5) (MORT1-ASSOCIATED CED-3 HOMOLOG) (MACH) (FADD-HOMOLOGOUS ICEJCED-3-LIKE PROTEASE) (FADD-LIKE ICE) (FLICE) (APOPTOTIC CYSTEINE PROTEASE) (APOPTOTIC PROTEASE MCH-5) (CAP4) (CASP8) (MCH5) isof | U60520; U58143; X98172; X98173; X98174; AF00962 | B3g | 1327-1607 |
| CASPASE-8 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 5) (MORT1-ASSOCIATED CED-3 HOMOLOG) (MACH) (FADD-HOMOLOGOUS ICE/CED-3-LIKE PROTEASE) (FADD-LIKE ICE) (FLICE) (APOPTOTIC CYSTEINE PROTEASE) (APOPTOTIC PROTEASE MCH-5) (CAP4) (CASP8) (MCH5) isof | $\begin{aligned} & \text { U60520; U58143; X98172; } \\ & \text { X98173; X98174; } \\ & \text { AF00962;X98176; X98175; } \\ & \text { X98177; X98178 } \end{aligned}$ | B3h | 475-954 |
| CASPASE-9 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 6) (ICE-LAP6) (APOPTOTIC PROTEASE MCH-6) | U56390; [U60521] | B3i | 986-1289 |
| ICE-LIKE APOPTOTIC PROTEASE 4 PRECURSOR (EC 3.4.22.-) (APOPTOTIC PROTEASE MCH-4) (CASPASE-10) | U60519 | B3j | 2276-2690 |
| DEATH-ASSOCIATED PROTEIN 3 (DAP-3) (ionizing radiation resistance conlerring protein) | U18321: [X83544] | B3K | 856-1114 |
| DEATH-ASSOCIATED PROTEIN KINASE 1 (EC 2.7.1.-) (DAP KINASE 1). | X76104 | B31 | 1988-2321 |


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| :---: | :---: | :---: | :---: |
| Fas-activated serine/threonine kinase (FAST) phosphorylates TIA-1 | X86779 | B3m | 865-1239 |
| PDCD2 | S78085 | B3n | 406-694 |
| FAS/APO 1 | 270519 | B4a | 1493-1887 |
| FAS ANTIGEN LIGAND (APOPTOSIS ANTIGEN LIGAND) (APTL) (APTILG1) (FASL). | D38122; [U08137] | B4b | 1400-1782 |
| WSL-LR, WSL-S1, WSL-S2 + TRAMP (Apo-3) (DDR3) | Y09392; [U75380;U74611; U83597] | B4c | 1407-1671 |
| Akt1 (rac protein kinase alpha, protein kinase B, c-Akt) | M63167 | B4d |  |
| AKT2 (rac protein kinase beta) | M77198; [M95936] | B4e | 1867-2099 |
| TNF-alpha converting enzyme | U69611 | B4I | 1540-1746 |
| death receptor 5 (DR5) | AF016268 | B4g | 273-552 |
| BRAG-1 = brain-related apoptosis gene/Bcl-2 homolog | S82185 | B4h | 351.995 |
| seven in absentia homolog | U63295 | B4i | 239-523 |
| RATS1 | U37688 | B4j | 1247-1367 |
| DNA fragmentation factor-45 | U91985 | B4k | 485-1592 |
| secreted apoplosis related protein 1 | AF017986 | B41 | 189-974 |
| secreted apoplosis related protein 3 (SARP3) | AF017988 | B4m | 702-841 |
| apoptosis-related protein TFAR15 (TFAR15) | AF022385 | B4n | 365.520 |
| calmodulin dependent phosphodiesterase PDE1B1 | U56976 | B5a | 414.549 |
| glutathione-S-transferase homolog | U90313 | B5b | 97-837 |
| CD27BP (Siva) | U82938 | B5c | 406-625 |
| chromosome segregation gene homolog CAS | U33286 | B5d | 674-1247 |
| apoptosis inhibitor survivin | U75285 | B5e | 386-720 |
| p53 induced protein | AF010310 AF010311 | B5I | 29.771 |
| Pig3 (PIG3) | AF010309 | B59 | 398-1223 |
| Pig7 (PIG7) | AF010312 | B5h | 173-322 |
| Pig10 (PIG10) | AF010314 | B5i | 437-1623 |
| Pig11 (PIG11) | AF010315 | B5 ${ }^{\text {B }}$ | 748-1304 |
| Pig12 (PIG12) | AF010316 | B5k | 97-531 |
| GTP-binding protein (rhoA) | L25080 | B5I | 290-572 |
| cdc42 homolog (G25K) [ brain isoform + placental isoform) | M35543; [M57298] | B5m | 321-468 |
| ONCOGENES/TUMOR SUPPRESSORS |  |  |  |
| C.FMS PROTO ONCOGENE | X03663 | B5n | 2568-2880 |
| C-los | K00650 | B6a | 2949-3181 |
| C-kit | X06182 | B6b | 1981-2375 |
| PROTO-ONCOGENE TYROSINE-PROTEIN KINASE SRC (EC 2.7.1.112) (P60-SRC) (C-SRC). | HT2291; [K03214; X03996] | B6c | 893-1189 |
| PROTO-ONCOGENE TYROSINE-PROTEIN KINASE FGR (EC 2.7.1.112) (P55-FGR) (C-FGR). | M19722 | B6d | 521-856 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| DNA MISMATCH REPAIR PROTEIN MSH2 | U04045; [L47583) | B6e | 1496-2178 |
| DNA MISMATCH REPAIR PROTEIN MSH6 (mutS - ALPHA 160 KD SUBUNIT) (G/T MISMATCH BINDING PROTEIN) (GTBP) (GTMBP) (P160) | U54777 | B6f | 591-1100 |
| K-RAS, ONCOGENE | M54968 | B6g | 352-604 |
| MET | J02958 | B6h | 932-1242 |
| p53 | M14694; [M14695] | B6i | 690-964 |
| BREAST CANCER TYPE 2 SUSCEPTIBILITY PROTEIN | U43746 | B6j | 10056-10346 |
| BRCAI-ASSOCIATED RING DOMAIN PROTEIN | U76638 | B6k | 1493-1801 |
| MDM2 PROTEIN (P53-ASSOCIATED PROTEIN) + MDM2-A (GB: U33199) + MDM2-C (GB: U33201) | Z12020; [M92424] | B61 | 920-1232 |
| MDM2-like p53-binding protein (MDMX) | AF007111 | B6m | 405-681 |
| p73, a monoallelically expressed p53-related protein | Y11416 | B6n | 627.993 |
| RB2/p130 | $\times 74594$ | B7a | 951-1213 |
| RBA/P48 | $\times 74262$ | B7b | 605-974 |
| RBP2 retinoblastoma binding protein | S66431 | B7c | 2339-2642 |
| RBQ1 retinoplastoma binding protein | X85133 | B7d | 1701-1930 |
| PROTO-ONCOGENE TYROSINE-PROTEIN KINASE RECEPTOR RET PRECURSOR (EC 2.7.1.112) (C-RET). [Papillary thyroid carcinomaencoded protein) | M31213; [M57464] | B7e | 2285-2631 |
| Retinoblastoma susceptibility (RB1 retinoblastoma-assoc) | M15400 | B71 | 2839-3101 |
| SKY (DTK) (TYRO3) (RSE) | D17517 | B7g | 2132-2597 |
| YES | M15990 | B7h | 1325-1676 |
| TYROSINE-PROTEIN KINASE BTK (EC 2.7.1.112) (BRUTON'S TYROSINE KINASE)(AGAMMAGLOBULINAEMIA TYROSINE KINASE) (ATK) (B CELL PROGENITOR KINASE) (BPK) (BTK) (AGMX1) | U10087 X58957 | B7i | 380-1430 |
| TYROSINE-PROTEIN KINASE ABL2 (EC 2.7.1.112) (TYROSINE KINASE ARG) (ABLL) | M35296 | B7j | 493-1656 |
| TYROSINE-PROTEIN KINASE ZAP-70 (EC 2.7.1.112) (70 KD ZETAASSOCIATED PROTEIN) (ZAP70) | L05148 | B7k | 1.584 |
| SIGNAL TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION 1ALPHABETA (TRANSCRIPTION FACTOR ISGF-3 COMPONENTS P91/P84) (STAT1) | M97935 | 871 | 638-1376 |
| SIGNAL TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION 2 (P113) (STAT2) | U18671 M97934 | B7m | 1105-1480 |
| SIGNAL TRANSDUCER AND TRANSCRIPTION ACTIVATOR 5B (STAT5B) | U47686 | B7n | $831-1135$ |
|  |  |  |  |
| QUADRANT C |  |  |  |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| DNA DAMAGE RESPONSEIREPAIR/RECOMBINATION |  |  |  |
| DNA -DEPENDENT PROTEIN KINASE (DNA-PK) + DNA DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT (DNA-PKcs) (XRCC7) | U35835; [U47077] | C1a | 2250-2680 |
| ATAXIA TELANGIECTASIA (ATM) | U33841 | C1b | 8938-9135 |
| FKBP.RAPAMYSIN ASSOCIATED PROTEIN (FRAP) | L34075 | C1c | 6750-7088 |
| ATP-DEPENDENT DNA HELICASE II, 70 KD SUBUNIT (LUPUS KU AUTOANTIGEN PROTEIN P70) (70 KD SUBUNIT OF KU ANTIGEN) (THYROID-LUPUS AUTO-ANTIGEN) (TLAA) (KU70) (CTC BOX BINDING FACTOR 75 KD SUBUNIT) (CTCBF) (CTC75) (XRCC6) | M32865 ; [S38729] | C1d | 1729-1974 |
| ATP-DEPENDENT DNA HELICASE II, 86 KD SUBUNIT (LUPUS KU AUTOANTIGEN PROTEIN P86) (86 KD SUBUNIT OF KU ANTIGEN) (THYROID-LUPUS AUTOANTIGEN) (TLAA) (CTC BOX BINDING FACTOR 85 KD SUBUNIT) (CTCBF) (CTC85) (NUCLEAR FACTOR IV) (KUBO) (XRCC5) | M30938 | C1e | 2340-2764 |
| DNA EXCISION REPAIR PROTEIN ERCC1 | M13194 | C11 | 625.938 |
| DNA LIGASE III (POLYDEOXYRIBONUCLEOTIDE SYNTHASE (ATP)) (DNL3) | $\times 84740$ | C1g | 2460-2780 |
| DNA LIGASE IV (POLYDEOXYRIBONUCLEOTIDE SYNTHASE (ATP)) (DNL4) | $\times 83441$ | C1h | 2787-3074 |
| DNA POLYMERASE ALPHA | X06745 | Cii | 3721-4093 |
| DNA REPAIR PROTEIN RAD50 | U63139 | C1i | 5117-5435 |
| DNA REPAIR PROTEIN RAD51 HOMOLOG [Replication protein A (E coli RecA homolog, RAD51 homolog)] | D13804 | C1k | 867-1159 |
| DNA REPAIR PROTEIN RAD52 HOMOLOG | U12134 | C11 | 1528-1733 |
| DNA TOPOISOMERASEI | J03250 | C1m | 2388-2796 |
| DNA TOPOISOMERASE II ALPHA ISOZYME | J04088 | C1n | 2459-2883 |
| DNA-REPAIR PROTEIN COMPLEMENTING XP-B CELLS (XERODERMA PIGMENTOSUM GROUP B COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROTEIN ERCC3) (BASAL TRANSCRIPTION FACTOR 289 KD SUBUNIT) (BTF2-p89) (TFIIH 89 KD SUBUNIT) | M31899 | C2a | 2109-2466 |
| DNA-REPAIR PROTEIN COMPLEMENTING XP.D CELLS (XERODERMA PIGMENTOSUM GROUP D COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROTEIN ERCC-2) | X52221; [HT1175] | C2b | 1520-1821 |
| DNA-REPAIR PROTEIN XRCC1 | M36089 | C2c | 1226-1539 |
| DNA-REPAIR PROTEIN COMPLEMENTING XP.G CELLS (XERODERMA PIGMENTOSUM GROUP G COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROTEIN ERCC-5) | L20046; [X69978 ] | C2d | 1374-1638 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| GROWTH ARREST AND DNA-DAMAGE-INDUCIBLE PROTEIN GADD153 (DNA-DAMAGE INDUCIBLE PROTEIN) (CHOP). | S40706 [S62138] | C2e | 480-789 |
| GROWTH ARREST AND DNA-DAMAGE-INDUCIBLE PROTEIN GADD45 (DNA-DAMAGE INDUCIBLE TRANSCRIPT 1) (DDIT1). | M60974 | C2f | 526-886 |
| METHYLATED-DNA--PROTEIN-CYSTEINE METHYLTRANSFERASE (6-O-METHYLGUANINE-DNA METHYLTRANSFERASE) (MGMT) | M29971 | C2g | 241-546 |
| MUSCLE-SPECIFIC DNASE I-LIKE [DNase X] (XIB) | X90392 ; [L40817; U06846] | C2h | 2038-2427 |
| DNA MISMATCH REPAIR PROTEIN MLH1 (mutL HOMOLOG) | U07418 | C2i | 1765-2020 |
| PAD | $L 24564$ | C2j | 489.780 |
| ACTIVATOR 136 KD SUBUNIT (REPLICATION FACTOR C 36 KD SUBUNIT) (RFC36) | L07540 | C2k | 708-1051 |
| ACTIVATOR 137 KD SUBUNIT (REPLICATION FACTOR C 37 KD SUBUNIT) (RFC37) | M87339 | C 21 | 98-355 |
| ACTIVATOR 138 KD SUBUNIT (REPLICATION FACTOR C 38 KD SUBUNIT) (RFC38) | L07541 | C2m | 438-762 |
| ACTIVATOR 140 KD SUBUNIT (REPLICATION FACTOR C 40 KD SUBUNIT) (RFC40) | M87338 | C2n | 882-1286 |
| REPLICATION PROTEIN A 70 KD DNA-BINDING SUBUNIT (RP-A) (RFA) (REPLICATION FACTOR-A PROTEIN 1) (SINGLE STRANDED DNABINDING PROTEIN) | M63488 | С3a | 1498-1838 |
| SUPEROXIDE DISMUTASE [Superoxide dismutase 1 (Cu/Zn)] | HT3218 [K00065] | C3b | 198-496 |
| TRANSCRIPTIONAL ACTIVATOR PROTEIN PUR-ALPHA | M96684 | C3c | 563-855 |
| HHR6A (YEAST RAD6 HOMOLOG) (UBIQITIN-CONJUGATING ENZYME) (UBCA) | M74524 | C3d | 175-433 |
| UV EXCISION REPAIR PROTEIN PROTEIN RAD23 [xeroderma pigmentosum group $C$ repair complementing protein HHR23A] | D21235 | C3e | 355-632 |
| CELL FATEJEVELOPMENT REGULATORS |  |  |  |
| - Notch pathway |  |  |  |
| Notch1 | M73980 | C3i | 2701-2965 |
| Notch2 | U77493 | C3g | 373-658 |
| notch group protein ( N ) | M99437 | C3h | 647-1210 |
| Notch4 | U95299 | C3i | 3014-3169 |
| Jagged 1 | AF028593 | C3j | 3884-4117 |
| Jagged 2 | AF003521 | C3k | 1027-1241 |
| DELTA-LIKE PROTEIN PRECURSOR (CONTAINS: FETAL ANTIGEN 1) <br> (FA1) (DLK) + ADRENAL SPECIFIC 30kd PROTEIN GB: X17544 | U15979; [Z12172] | C31 | 1090-1403 |
| manic fringe | U94352 | C3m | 979-1235 |



| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| IGFBP3 (GROWTH HORMONE-DEPENDENT INSULIN-LIKE GROWTH FACTOR-BINDING PROTEIN) | M31159; [M35878] | C6d | 451.744 |
| IGFBP4 | M62403 | C6e | 657.967 |
| IGFBP5 | M65062 | C6! | 356-602 |
| IGFBP6 | M62402 | C69 | 345-536 |
| INSULIN-LIKE GROWTH FACTOR I RECEPTOR | X04434 | C6h | 3413-3904 |
| BASIC FIBROBLAST GROWTH FACTOR RECEPTOR 1 PRECURSOR (BFGF-R) (EC 2.7.1.112) (FMS-LIKE TYROSINE KINASE-2) (C-FGR) (FGFR1) (FLG) (FGFBR) (FLT2). (HBGF-R-ALPHA-A1) (HBGF-R-ALPHAA2) (HBGF-R-ALPHA-A3) + FGFR SECRETED FORM (M34188) | M37722; [X66945; M63887; M63888; M63889;M34186; M34641] | C6i | 1746-1967 |
| NERVE GROWTH FACTOR RECEPTOR | M14764 | C6j | 2762-3242 |
| PDGFR-ALPHA | M21574 | C6k | 5118-5583 |
| PDGFR-BETA | M21616 | C61 | 842-1133 |
| transmembrane receptor precursor (PTK7); COLON CARCINOMA KINASE-4 (CCK4) | U33635; [U40271] | C6m | 3507-3784 |
| SEX GENE | X87852 | C6n | 209-433 |
| TRANSFORMING GROWTH FACTOR-BETA TYPE III RECEPTOR | L07594 | C7a | 3358-3592 |
| TRANSMEMBRANE PROTEIN TMP21 | X97442 | C7b | 380-1176 |
| HIGH AFFINITY NERVE GROWTH FACTOR RECEPTOR PRECURSOR (EC 2.7.1.112) (TRK1 TRANSFORMING TYROSINE KINASE PROTEIN) (P140-TRKA) + trk-T3 (P68 TRK-T3 ONCOPROTEIN) | X03541 | C7c | 1816-2118 |
| irk-T3 (P68 TRK-T3 ONCOPROTEIN) | X85960 | C7d | 252-1112 |
| trk-B | U12140 | C7e | 1006-1384 |
| trk-C | U05012 | C7t | 359.765 |
| TUMOR NECROSIS FACTOR RECEPTOR 1 | M33294 | C7g | 1570-1817 |
| TUMOR NECROSIS FACTOR RECEPTOR 2 PRECURSOR (TUMOR NECROSIS FACTOR BINDING PROTEIN 2) (TBPII) (P80) (TNF-R2) (P75) (CD120B) (TNFR2) (TNFBR). | M32315; [M55994] | C7h | 3359-3543 |
| RETINOIC ACID RECEPTOR ALPHA1 (RAR- ALPHA1) + PML-RAR protein | $\begin{aligned} & \text { M73779; [X06538; } \\ & \text { (X06614] } \end{aligned}$ | C7i | 2935-3238 |
| retinoic acid receptor alpha [RETINOIC ACID RECEPTOR RXR-ALPHA (RXRA)] | X52773 | C7i | 352-616 |
| retinoic acid receptor epsilon [RETINOIC ACID RECEPTOR BETA-2 (RAR BETA-2) (RAR-EPSILON)I | X07282; [Y00291] | C7k | 1315-1633 |
| retinoic acid receptor gamma [RETINOIC ACID RECEPTOR GAMMA] | $\begin{aligned} & \text { M24857; [M38258; } \\ & \text { M57707; M32074] } \end{aligned}$ | C71 | 1569-1834 |
| retinoic acid receptor rxr-beta [RETINOIC ACID RECEPTOR RXR-BETA] | M84820; [X63522] | C7m | 643-1135 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| THROMBOPOEITIN RECEPTOR | U68162 | C7n | 5117-5435 |
| QUADRANT D |  |  |  |
| CELL ADHESION, MOTILITY, AND INVASION |  |  |  |
| CARTILAGE-SPECIFIC PROTEOGLYCAN CORE PROTEIN (CSPCP) (AGGRECAN 1)(CHONDROITIN SULFATE PROTEOGLYCAN CORE PROTEIN 1) | M55172 | D1a | 6705-6956 |
| byglycan | J04599 | D1b | 854-1129 |
| CD34 | M81104 | D1c | 596-960 |
| CD59 | M34671 | D1d | 105-1163 |
| CHONDROITIN/DERMATAN SULFATE PROTEOGLYCAN CORE PROTEIN (DECORIN) (PG-S2) (PG40) | M14219 | D1e | 712-896 |
| COLLAGEN (-6000BP) | D21337 | D11 | 5342-5588 |
| collagen type l | X55525 | D19 | 428-741 |
| collagen type II alpha-1 | X16468 | D1h | 3604-3751 |
| collagen type III pro-alpha-1 | X14420 | D1i | 3867-4046 |
| collagen type IV alpha | X05610 | D1i | 882-1113 |
| collagen type IV alpha-3 | M92993 | D1k | 2296-2545 |
| collagen type VI alpha-1 | X15879 | D11 | 316-688 |
| collagen type VI alpha-2 | M34570 | D1m | 203-396 |
| collagen type VI alpha-3 | X52022 | D1n | 640-1487 |
| collagen type Vill alpha- 1 | $\times 57527$ | D2a | 612.1772 |
| collagen type XI alpha-1 | $J 04177$ | D2b | 2864-3091 |
| collagen type XI pro-alpha-2 | U32169 | D2c | 4473-4769 |
| collagen type XVI alpha-1 | M92642 | D2d | 4816-5991 |
| collagen type XVIII alpha | L22548 | D2e | $2300-2539$ |
| LAM3AH (LAMA4) | X70904; [ $\times 91171$ ] | D21 | 1018-1388 |
| LAMB2 (LAMININ) | 577512 | D2g | $3871-4158$ |
| laminin B1 | M61916 | D2h | 3177-3554 |
| laminin B2 | J03202 | D2i | 2878-3232 |
| laminin, 37KD RECEPTOR | U43901 | D2j | 460-812 |
| netrin-2 | U86759 | D2k | 859-1147 |
| nidogen | M30269 | D21 | 2120-2428 |
| TENASCIN-C | X78565 | D2m | 6652-6924 |
| TENASCIN-R | X98085 | D2n | 3916-4165 |
| VERSICAN [isoforms . V1, V2, V3] | $\begin{aligned} & \text { U16306; [X15998; U26555; } \\ & \text { D32039] } \end{aligned}$ | D3a | 189-974 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| SPARC PRECURSOR (SECRETED PROTEIN ACIDIC AND RICHIN CYSTEINE) (OSTEONECTIN) (ON) (BASEMENT MEMBRANE PROTEIN BM-40). | J03040 | D3b | 280.642 |
| THROMBOSPONDIN 1 PRECURSOR | X14787 | D3c | 3187-3450 |
| THROMBOSPONDIN 2 PRECURSOR | L12350 | D3d | 3151-3531 |
| VITRONECTIN PRECURSOR (SERUM SPREADING FACTOR) (SPROTEIN) (CONTAINS: SOMATOMEDIN B) | X03168 | D3e | 3721-4093 |
| fibronectin | $\times 02761$ | D3i | 6163-7290 |
| RNA-binding protein Hel-N2; ELAV-like neuronal protein 1 | U12431; [U29943] | D39 | 1006-1384 |
| HEPARAN SULFATE PROTEOGLYCAN (HSPG2) | M85289 | D3h | 1232-1389 |
| integrin alpha | X68742 | D3i | 2690-2976 |
| integrin alpha2 [very late antigen-2 (vla-2)/collagen receptor alpha-2 subunit] | M28249; [X17033] | D3j | 2367-2664 |
| integrin alpha3 | M59911 | D3k | 2564-2944 |
| integrin alpha4 | L12002; [ $\times 16983$ ] | D31 | 2709-3063 |
| integrin alpha5 [fibronectin receptor alpha subunit] | X06256 | D3m | 2094-2367 |
| integrin alpha6 | X53586; [X59512] | D3n | 3642-3988 |
| integrin alpha7B | X74295 | D4a | 255-591 |
| integrin alpha8 | L36531 | D4b | 2709-3063 |
| integrin alpha9 | D25303; [L24158] | D4c | 706-980 |
| integrin alphaE | L25851 | D4d | 2279-2529 |
| integrin betal | M34189 | D4e | 701-1301 |
| integrin beta3 [PLATELET MEMBRANE GLYCOPROTEIN IIIA] | J02703; [M25108] | D4f | 2038-2373 |
| inlegrin beta 4 | X53587; [X52186] | D49 | 5357-5697 |
| integrin beta5 | J05633 | D4h | 2279-2528 |
| integrin beta6 | M35198 | D4i | 1619-1901 |
| integrin beta7 | M62880 | D4j | 2562-2944 |
| integrin beta8 | M73780 | D4k | 22-877 |
| Focal adhesion kinase | L13616 | D41 | 2179-2631 |
| Integrin-linked kinase (ILK) | U40282 | D4m | 1245-1530 |
| Protein tyrosine kinase Pyk2 (Cell adhesion kinase-beta, CAK-beta) (FAK2) | U43522; [L49207] | D4n | 3658-3952 |
| Paxillin | U14588 | D5a | 1260-1644 |
| Zyxin + Zyxin-2 | X94991; [X95735] | D5b | 585-1514 |
| Zyxin related protein ZRP-1 | AF000974 | D5c | 1240-1466 |
| beta 3-endonexin | U37139 | D5d | 606.1504 |
| cytohesin-1; Sec7p-like protein | U59752 | D5e | 43-338 |
| CD9 | M38690 | D5f | 372.962 |
| Ezrin (cytovillin 2) | $\times 51521$ | D5g | 1611-1883 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| MERLIN (SCHWANNOMIN) (moesin-ezrin-radixin-like protein)(neurofibromatosis 2) | $\begin{aligned} & \text { L11353; Z22664; X72657; } \\ & \text { L27133 } \end{aligned}$ | D5h | 355-674 |
| LICAM | M74387 | DSi | 3197-3485 |
| N.CAM [NEURAL CELL ADHESION MOLECULE, PHOSPHATIDYLINOSITOL-LINKED ISOFORM; CD56] | X16841 | D5j | 2338-2646 |
| NINJURIN-1 | 072661 | D5k | 212-492 |
| opioid binding cell adhesion molecule | L34774 | D51 | 115.728 |
| DCC | X76132 | D5m | 893-1189 |
| P37NB | U32907 | D5n | 95-456 |
| PLEXIN | U52111 | D6a | 585-1514 |
| semaphorin (CD100) | U60800 | D6b | 2517-2921 |
| semaphorin E | AB000220 | D6c | 2949-3181 |
| semaphorin III | $\underline{L 26081}$ | D6d | 899-1152 |
| semaphorin V | U33920 | D6e | 177-442 |
| SEMAPHORIN-1 | U38276 | D6f | 488-653 |
| TAX1, AXONIN-1/TAQ1 | $\times 85978$ | D69 | 209-433 |
| LAR | Y00815 | D6h | 5799.6049 |
| HYALURONAN RECEPTOR (RHAMM) | U29343 | D6i | 2496-2798 |
| PLATELET GLYCOPROTEIN IV (GPIV) (GPIIIB) (CD36 ANTIGEN) (PAS IV) (PAS-4 PROTEIN) | M24795 | D6j | 554-806 |
| caveolin-2 | AF035752 U32114 | D6k | 1340-1519 |
| caveolin-1 | Z18951 249856 | D61 | 62-413 |
| ANGIOGENESIS REGULATORS |  |  |  |
| VASCULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR 2 PRECURSOR (EC 2.7.1.112) (VEGFR-2) (KDR) (KINASE INSERT DOMAIN RECEPTOR) (FRAGMENT) | L04947; [X61656] | D6m | 2686-3053 |
| VASCULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR 3 PRECURSOR (EC 2.7.1.112) (VEGFR-3) (TYROSINE-PROTEIN KINASE RECEPTOR FLT4, CLASS III). | X68203; [X69878; U43143] | D6n | 4236-4402 |
| FL CYTOKINE RECEPTOR PRECURSOR (EC 2.7.1.112) (TYROSINEPROTEIN KINASE RECEPTOR FLT3) (STEM CELL TYROSINE KINASE 1) (STK-1) (CD135 ANTIGEN). | U02687 | D7a | 2491-2965 |
| TYROSINE-PROTEIN KINASE RECEPTOR TIE-1 PRECURSOR (EC 2.7.1.112). | X60957 [S89716] | D7b | 3114-3536 |
| TYROSINE-PROTEIN KINASE RECEPTOR TIE-2 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR TEK) (P140 TEK) ( (TUNICA INTERNA ENDOTHELIAL CELL KINASE). | L06139 | D7c | 3243-3586 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| VASCULAR ENDOTHELIAL GROWTH FACTOR B PRECURSOR (VEGFB) + VEGF RELATED FACTOR ISOFORM VRF186 PRECURSOR | U48801; [U43368] | D7d | 158-648 |
| VASCULAR ENDOTHELIAL GROWTH FACTOR C PRECURSOR (VEGF- <br> C) (VASCULAR ENDOTHELIAL GROWTH FACTOR RELATED PROTEIN) (VRP) (FLT4 LIGAND). | U43142 | D7e | 1165-1559 |
| PLACENTA GROWTH FACTORS 1 AND 2 PRECURSOR (PLGF-1/ PLGF-2). | $\times 54936$ | D71 | 1098-1371 |
| SL CYTOKINE PRECURSOR (FLT3/FLK2 LIGAND). | U04806; [U03858] | D79 | 29-362 |
| angiopoietin-1 | U83508 | D7h | 1749-2031 |
| CYSTEINE-RICH FIBROBLAST GROWTH FACTOR RECEPTOR [Golgi membrane sialoglycoprotein MG160 (GLG1)] | U28811; [U64791] | D7i | 3279-4140 |
| FGFR3 (FLG-2) | M58051; [X58255] | D7j | 323-896 |
| FGFR4 | L03840 | D7k | 1503-1743 |
| FIBROBLAST GROWTH FACTOR RECEPTOR 2 PRECURSOR (FGFR2) (EC 2.7.1.112) (KERATINOCYTE GROWTH FACTOR RECEPTOR) (FGFR2) (BEK) (BFR-1) (KSAM-1) + K-SAM; K-SAM-III; K-SAM-IV | U11814; [M80634; X52832; M35718; M87771; M87772] | D71 | 753-1189 |
| VASCULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR 1 PRECURSOR (EC 2.7.1.112) (VEGFR-1) (TYROSINE-PROTEIN KINASE RECEPTOR FLT) (FLT-1) (SFLT) | U01134; [X51602] | D7m | 1288-1604 |
| HOMEOBOX PROTEIN HOX-D3 [HOX 4A] | D11117 | D7n | 4200-4447 |
| QUADRANT E |  |  |  |
| INVASION REGULATORS |  |  |  |
| MMP-1 (collagenase-1) | X05231 | E1a | 512-836 |
| MMP-2 (gelatinase A) | J03210, [J05471] | E1b | 477.778 |
| MMP-3 (stromelysin-1) | $\times 05232$ | E1c | 331-1491 |
| MMP-7 (matrilysin) | $\times 07819$ | E1d | 335-738 |
| MMP-8 (collagenase-2) | J05556 | E1e | 532.865 |
| MMP-9 (gelatinase B) | J05070, [D10051] | E1f | 1012-1346 |
| MMP-10 (stromelysin-2) | X07820, [M30461] | E1g | 387-1319 |
| MMP-11 (stromelysin-3) | X57766 | E1h | 263-1508 |
| MMP-12 (metalloelastase) | L23808 | E1i | 275-787 |
| MMP-13 (collagenase-3) | X75308 | E1j | 463.761 |
| MMP-14 (MT1-MMP) | D26512, [ $\times 83535$ ] | E1k | 413-749 |
| MMP-15 (MT2-MMP) | 248482 | E11 | 1210-1456 |
| MMP-16 (MT3-MMP) | D50477 | E1m | 991-1226 |
| MMP. 17 (MT4-MMP) | X89576 | E1n | 630-1830 |
| MMP-19 | $\times 92521$ | E2a | 1383-1655 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| TIMP-1 (erythroid polentiating activity, EPA) | X03124 | E2b | 194.492 |
| TIMP-2 (MI) | $J 05593$ | E2c | 403-694 |
| TIMP-3 (mitogen-inducible gene 5, mig-5) | Z30183 | E2d | 346-587 |
| TIMP-4 | U76456 | E2e | 445-671 |
| extracellular matrix metalloproteinase inducer EMMPRIN | L20471 | E2f | 23-354 |
| UROKINASE-TYPE PLASMINOGEN ACTIVATOR PRECURSOR (EC 3.4.21.73) (UPA) (U-PLASMINOGEN ACTIVATOR) | M15476 | E2g | 824-1120 |
| TISSUE-TYPE PLASMINOGEN ACTIVATOR PRECURSOR (EC 3.4.21.68) (T-PA) (T-PLASMINOGEN ACTIVATOR). | $\begin{aligned} & \begin{array}{l} \text { M15518; [ X07393; } \\ \text { M18182) } \end{array} \\ & \hline \end{aligned}$ | E2h | 1221-1577 |
| PLASMINOGEN PRECURSOR (EC 3.4.21.7) | $\times 05199$ | E2i | 1859-2162 |
| PLASMINOGEN ACTIVATOR INHIBITOR-1 PRECURSOR, ENDOTHELIAL (PAI-1) | $\times 04429$ | E2j | 1195-1342 |
| PLASMINOGEN ACTIVATOR INHIBITOR-2, PLACENTAL (PAI-2) (MONOCYTE ARG- SERPIN) (UROKINASE INHIBITOR). | M18082; J02685] | E2k | 378-954 |
| PLASMA SERINE PROTEASE INHIBITOR PRECURSOR (PCI) (PROTEIN C INHIBITOR) (PLASMINOGEN ACTIVATOR INHIBITOR-3) (PA13). | M68516; [J02639] | E21 | 8035-8423 |
| UROKINASE PLASMINOGEN ACTIVATOR SURFACE RECEPTOR, GPIANCHORED FORM PRECURSOR (U-PAR) (MONOCYTE ACTIVATION ANTIGEN MO3) (CD87 ANTIGEN) | U08839 [M83246; X51675] | E2m | 749-1043 |
| LOW-DENSITY LIPOPROTEIN RECEPTOR-RELATED PROTEIN 1 PRECURSOR (LRP) (ALPHA-2-MACROGLOBULIN RECEPTOR) (A2MR) | X13916 | E2n | 5439-5742 |
| LOW-DENSITY LIPOPROTEIN RECEPTOR-RELATED PROTEIN 2 (MEGALIN) (GLYCOPROTEIN 330) (FRAGMENT) | U04441 | E3a | 1365-2162 |
| ALPHA-2-MACROGLOBULIN PRECURSOR (ALPHA-2-M) | M11313 | E3b | 3972-4325 |
| PLATELET BASIC PROTEIN PRECURSOR (PBP) (CONTAINS: CONNECTIVE-TISSUE ACTIVATING PEPTIDE III (CTAP-III), LOWAFFINITY PLATELET FACTOR IV (LA-PF4), BETA- <br> THROMBOGLOBULIN (BETA-TG), NEUTROPHIL-ACTIVATING PEPTIDE 2 (NAP-2)) | M54995; M38441 | E3c | 63-252 |
| ALPHA-2-MACROGLOBULIN RECEPTOR-ASSOCIATED PROTEIN PRECURSOR (ALPHA-2-MRAP) (LOW DENSITY LIPOPROTEIN RECEPTOR-RELATED PROTEIN- ASSOCIATED PROTEIN 1) (RAP) | M63959 | E3d | 440-890 |
| NUCLEOSIDE DIPHOSPHATE KINASE A (EC 2.7.4.6) (NDK A) (NDP KINASE A) (TUMOR METASTATIC PROCESS-ASSOCIATED PROTEIN) (METASTASIS INHIBITION FACTOR NM23) (NM23-H1). | X17620 | E3e | 245-612 |



| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| CADHERIN-4 RETINAL-CADHERIN PRECURSOR (R-CADHERIN) (RCAD) | L34059 | ESg | 1172-1425 |
| CADHERIN-5 VASCULAR ENDOTHELIAL-CADHERIN PRECURSOR (VE CADHERIN) (7B4 ANTIGEN) (CD144 ANTIGEN). | X79981; [X59796] | E5h | 1607-1769 |
| CADHERIN-6 | D31784 | E5i | 2119.2443 |
| CADHERIN-8 | L34060 | E5i | 1069-1347 |
| CADHERIN-11 (OSTEOBLAST-CADHERIN) (OB-CADHERIN) | L34056 | E5k | 1778-2076 |
| CADHERIN-12 (BR-CADHERIN) (N-CADHERIN 2) (CADHERIN, NEURAL TYPE, 2) | L34057; [L33477] | E5I | 657.903 |
| CADHERIN-13 T-CADHERIN PRECURSOR (TRUNCATED-CADHERIN) (H-CADHERIN) (HEART-CADHERIN) | L34058; [U59289; U59288] | E5m | 949-1187 |
| CADHERIN-14 MUSCLE-CADHERIN PRECURSOR (M-CADHERIN) (CADHERIN-14) (CADHERIN-15) | D83542 | E5n | 228-456 |
| ALPHA-CATENIN (CADHERIN-ASSOCIATED PROTEIN) (ALPHA E- CATENIN) | D13866 [D14705 L23805; L22080] | E6a | 55-492 |
| ALPHA-CATENIN RELATED PROTEIN (CATENIN ALPHA-2) | M94151 | E6b | 2296-2545 |
| BETA-CATENIN | X87838 [219054] | E6c | 2061-2463 |
| PLAKOGLOBIN (DESMOPLAKIN III) | M23410 | E6d | 2000-2312 |
| APC (DP2.5) | M74088; (M73548) | E6e | 7992-8326 |
| neuroendocrine-dig (NE-dig) a novel human homolog of the Drosophila discs large (dig) tumor suppressor protein interacting with the APC protein | U49089 | E6 | 2210-3116 |
| EB1, a protein that binds to APC | U24166 | E6g | 488.796 |
| protocadherin 42 | L11370 | E6h | 1246-1605 |
| protocadherin 43 | L11373 | E6i | 1018-1388 |
| desmoplakin 1 | M77830 | E6j | 6987-7826 |
| envoplakin (EVPL) | U53786 | E6k | 5583.5788 |
| bullous pemphigoid antigen | M63618 | E61 | 5680-6055 |
| desmoglein 2 | Z26317 [S64273] | E6m | 2819-3135 |
| desmoglein type 1 | $\times 56654$ | E6n | 2578-2889 |
| desmocollin type 1 | $\times 72925$ | E7a | 475-1154 |
| desmocollin type $3+$ desmocollin type 4 | X83929; (D17427) | E7b | 608-1607 |
| DSC2 mRNA for desmocollins type 2a and 2b | $\times 56807$ | E7c | 802-1115 |
| EPHRIN-A1 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 1) (LERK-1) (IMMEDIATE EARLY RESPONSE PROTEIN B61) (TUMOR NECROSIS FACTOR, ALPHA-INDUCED PROTEIN 4). | M57730 M37476 | E7d | 124-1062 |
| EPHRIN-A5 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 7) (LERK-7) (AL-1). | U26403 | E7e | 375-1325 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| EPHRIN-B1 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 2) (LERK-2) (ELK LIGAND PRECURSOR) (ELK-L). | U09304 | E7f | 507-1186 |
| EPHRIN-B2 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 5) (LERK-5) (HTK LIGAND) (HTK-L). | L38734 | E79 | 442-560 |
| EPHRIN-B3 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 8) (LERK-8) (EPH-RELATED RECEPTOR TRANSMEMBRANE LIGAND ELK-L3). | U66406 | E7h | 2056-2282 |
| EPHRIN TYPE-A RECEPTOR 2 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR ECK) (EPITHELIAL CELL KINASE). | M59371 M36395 | E7i | 249-1426 |
| EPHRIN TYPE-A RECEPTOR 5 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR EHK-1) (EPH HOMOLOGY KINASE-1) (RECEPTOR PROTEIN- TYROSINE KINASE HEK7). | X95425 | E7j | 644-1300 |
| EPHRIN TYPE-B RECEPTOR 1 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR EPH-2) (NET). | L40636 | E7k | 998-1469 |
| EPHRIN TYPE-B RECEPTOR 2 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN EPH-3) (DRT) | L41939 | E71 | 454-1225 |
| EPHRIN TYPE-B RECEPTOR 4 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR HTK). | U07695 | E7m | 756-1652 |
| TYROSINE-PROTEIN KINASE HCK (EC 2.7.1.112) (P59-HCK AND P60HCK) (HEMOPOIETIC CELL KINASE). | M16591 | E7n | 194-1187 |
|  |  |  |  |
| QUADRANT F |  |  |  |
| GROWTH FACTORS/CYTOKINES |  |  |  |
| AMPHIREGULIN | M30704 | F1a | 511-837 |
| BCGF1 (B-cell growth factor) | M15530 | F1b | 13.248 |
| BDNF | M61176 | F1c | 982-1265 |
| BETA NGF | X52599 | F1d | 360-1339 |
| VASCULAR ENDOTHELIAL GROWTH FACTOR PRECURSOR (VEGF) (VASCULAR PERMEABILITY FACTOR) (VPF). | M32977: [M27281] | F1e | 198-622 |
| BIGH3 | M77349 | F1f | 705-1703 |
| BONE MORPHOGENETIC PROTEIN 1 (procollagen C-proteinase) (pCP2) | M22488; [U50330] | F19 | 702-1098 |
| BONE MORPHOGENETIC PROTEIN 2A | M22489 | F1h | 567-997 |
| BONE MORPHOGENETIC PROTEIN 3 | M22491 | F1i | 1458-1731 |
| BONE MORPHOGENETIC PROTEIN 3B | D49493 | F1j | 16188-16418 |
| BONE MORPHOGENETIC PROTEIN 4 (BMP-2B) | D30751: [M22490] | Fik | 943-1321 |
| BONE MORPHOGENETIC PROTEIN 5 | M60314 | F11 | 1679-1982 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| BONE MORPHOGENETIC PROTEIN 6 | M60315 | F1m | 1067.1327 |
| BONE MORPHOGENETIC PROTEIN 7 (OSTEOGENIC PROTEIN 1) | M60316 | Fin | 451-691 |
| BONE MORPHOGENETIC PROTEIN 8 (OSTEOGENIC PROTEIN 2) | M97016 | F2a | 1345-1645 |
| BPGF- 1 | L42379 | F2b | 825-1213 |
| CNTF, ISOFORM B AND C | A26792 | F2c | 213-448 |
| CONNECTIVE TISSUE GROWTH FACTOR | M92934 | F2d | 1459-1748 |
| EGF (kidney) | $\times 04571$ | F2e | 4164-4434 |
| EGF-LIKE GROWTH FACTOR | M60278 | F21 | 1905-2146 |
| endothelin 2 | M65199 | F2g | 338-570 |
| endothelin 3 | J05081 | F2h | 1428-1685 |
| HEPARIN-BINDING GROWTH FACTOR 1 PRECURSOR (HBGF-1) (ACIDIC FIBROBLAST GROWTH FACTOR) (AFGF) (BETAENDOTHELIAL CELL GROWTH FACTOR) (ECGF-BETA). | X51943; [M13361; X65778] | F2i | 1131-1502 |
| FGF2; HEPARIN-BINDING GROWTH FACTOR 2 PRECURSOR (PROSTATROPIN). (HBGF-2) (BASIC FIBROBLAST GROWTH FACTOR) (BFGF) (PROSTATROPIN) | M27968 | F2j | 1384-1646 |
| FGF-3: INT-2 PROTO-ONCOGENE PROTEIN PRECURSOR (FIBROBLAST GROWTH FACTOR-3)(HBGF-3). | X14445 | F2k | 189-940 |
| FGF-5; FIBROBLAST GROWTH FACTOR-5 PRECURSOR (HBGF-5). | M37825 | F21 | 603-1086 |
| FGF-6; FIBROBLAST GROWTH FACTOR-6 PRECURSOR (HBGF-6) | X63454 | F2m | 287-456 |
| FGF-7; KERATINOCYTE GROWTH FACTOR PRECURSOR (KGF) (FIBROBLAST GROWTH FACTOR- 7) (HBGF-7). | M60828 | F2n | 522.955 |
| FGF-8; ANDROGEN-INDUCED GROWTH FACTOR PRECURSOR (AIGF) (HBGF-8) (FIBROBLAST GROWTH FACTOR-8) | U36223 | F3a | 32-3106 |
| FGF-9; GLIA-ACTIVATING FACTOR PRECURSOR (GAF) (FIBROBLAST GROWTH FACTOR-9) (HBGF-9). | D14838 | F3b | 110-949 |
| FHF-1 | U66197 | F3c | 17-566 |
| GDNF | L19063 | F3d | 248-390 |
| GLIA MATURATION FACTOR beta | $\begin{aligned} & \text { HG563 [M86492; } \\ & \text { AB001106] } \end{aligned}$ | F3e | 203-434 |
| RECOMBINANT GLIAL GROWTH FACTOR + NEU DIFFERENTIATION FACTOR + HEREGULIN | L12260; U02326; M94165 | F3i | 1069-1452 |
| TRANSFORMING GROWTH FACTOR-BETA-2 (glioblastoma-derived tcell suppressor factor) | M19154; [Y00083] | F39 | 1538-1878 |
| GROWTH INHIBITORY FACTOR (METALLOTHIONEIN-III) (MT-III) | D13365: [M93311] | F3h | 4-1052 |


| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| PLEIOTROPHIN PRECURSOR (PTN) (HEPARIN-BINDING GROWTHASSOCIATED MOLECULE) (HB-GAM) (HEPARIN-BINDING GROWTH FACTOR 8) (HBGF-8) (OSTEOBLAST SPECIFIC FACTOR 1) (OSF-1) (HEPARIN-BINDING NEURITE OUTGROWTH PROMOTING FACTOR 1) (HBNF-1). | M57399; [X52946; D90226] | F3i | 602-847 |
| EARLY GROWTH RESPONSE PROTEIN 1 (EGR-1) (KROX24) (TRANSCRIPTION FACTOR ETR103) (ZINC FINGER PROTEIN 225) (AT225). | M62829; [X52541] | F3j | 989-1276 |
| HEPATOCYTE GROWTH FACTOR-LIKE (macrophage-stimulating protein (MST1)) | M74178 | F3k | 1643-2015 |
| HEPTOMA-DERIVED GROWTH FACTOR | D16431 | F31 | 359-625 |
| HEPATOCYTE GROWTH FACTOR PRECURSOR (SCATTER FACTOR) (SF) (HEPATOPOEITIN A). | M60718 | F3m | 1549-1970 |
| HGF AGONIST/ANTAGOINST | U46010 | F3n | 895-1051 |
| COMPETITIVE HEPATOCYTE GROWTH FACTOR ANTAGONIST. AN ALTERNATIVE TRANSCRIPT OF THE HEPATOCYTE GROWTH FACTOR PRECURSOR (SCATTER FACTOR) (SF) (HEPATOPOEITIN A) | M77227 | F4a | 947-1968 |
| IFN-GAMMA ANTAGONIST CYTOKINE | A25270 | F4b | 395-685 |
| IGF.1 | M27544; [M37484] | F4c | 652-919 |
| INTERLEUKIN 1 RECEPTOR ANTAGONIST | M63099 | F4d | 225-1294 |
| INTERLEUKIN 6 RECEPTOR | M20566 | F4e | 2359-2823 |
| INTERLEUKIN IL-1 ALPHA | X02851 | F4if | 1107-1473 |
| INTERLEUKIN IL-1BETA | K02770 | F4g | 917-1208 |
| INTERLEUKIN IL-2 | A14844 | F4h | 181.436 |
| INTERLEUKIN-3 PRECURSOR (IL-3) (MULTIPOTENTIAL COLONYSTIMULATING FACTOR) (HEMATOPOIETIC GROWTH FACTOR) (PCELL STIMULATING FACTOR) (MAST-CELL GROWTH FACTOR) (MCGF) (IL3). | M14743; [M17115] | F4i | 390-608 |
| INTERLEUKIN IL-4 | M13982 | F4j | 216-459 |
| INTERLEUKIN IL-5 (B CELL DIFFERENTIATION FACTOR I) (T-CELL REPLACING FACTOR) (EOSINOPHIL DIFFERENTIATION FACTOR) | X04688; [J03478] | F4k | 35-279 |
| INTERLEUKIN-6 PRECURSOR (IL-6) (B-CELL STIMULATORY FACTOR 2) (BSF-2) (INTERFERON BETA-2) (HYBRIDOMA GROWTH FACTOR). | X04602; [M14584] | F41 | 130-555 |
| INTERLEUKIN IL-7 | J04156 | F4m | 174-447 |
| INTERLEUKIN IL-9 (P40) | X17543; (M30134] | F4n | 156-399 |
| INTERLEUKIN IL-10 | M57627 | F5a | 442.648 |
| INTERLEUKIN IL-11 [adipogenesis inhibitory factor] | M57765 | F5b | 132.460 |
| INTERLEUKIN IL-12 (NKSF, P35) | M65291 | F5c | 600-990 |



| Cell Cycle/Growth Regulators | GenBank \# | Array Coordinate | Position |
| :---: | :---: | :---: | :---: |
| INTERFERON-GAMMA RECEPTOR BETA CHAIN [Interferon gamma receptor accessory factor-1 (AF-1)] | U05875 | F71 | 1702-2039 |
| INTERFERON REGULATORY FACTOR [Interferon regulatory factor 1] | X14454 | F7g | 478-695 |
| INTERFERON CONSENSUS SEQUENCE BINDING PROTEIN [DNA- binding protein] | M91196 | F7h | 1253-1475 |
| HUIFN- ALPHA-REC [INTERFERON ALPHA-BETA RECEPTOR ALPHA CHAIN] | J03171 | F7i | 2562-2740 |
| INTERFERON ALPHA-BETA RECEPTOR BETA CHAIN | X77722 | F7i | 553-1012 |
| INTERFERON-GAMMA RECEPTOR ALPHA CHAIN | J03143 | F7k | 610-824 |
| INTERFERON-GAMMA RECEPTOR | A09781 | F71 | 66-317 |
| GAMMA INTERFERON INDUCED MONOKINE [Humig] | $\times 72755$ | F7m | 2021-2246 |
| INTERFERON-GAMMA INDUCED PROTEIN | $\times 02530$ | F7n | 280-613 |
| HOUSEKEEPING GENES |  |  |  |
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## Apoptosis Array

In the apoptosis array according to the subject invention, all of the unique polynucleotide probe compositions correspond to genes that are associated with apoptosis, e.g. cell cycle genes. In a specific apoptosis array of interest, the spots are as provided in Table 4.


|  | GenBank\# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: | :---: |
|  | U47414 [L49506] | CYCLIN 22 | 5 D |  |
|  | U11791 [U12685] | CYCLINH | 5 5 |  |
|  | U09579; [L25610] | CYCLIN-DEPENDENT KINASE INHIBITOR 1 (MELANOMA DIFFERENTIATION ASSOCIATED PROTEIN 6) (MDA-6) (P21) (CDK-INTERACTING PROTEIN 1) (CIP1) (WAF1) (CDKN1A) (CDKN1) (SDI1) (PIC1) (CAP20) | 5 F |  |
|  | U22398 | CYCLIN-DEPENDENT KINASE INHIBITOR 1C (CYCLIN-DEPENDENT KINASE INHIBITOR P57) (P57KIP2) | 5 G |  |
|  | L27211 | CYCLIN-DEPENDENT KINASE 4 INHIBITOR A (CDK4I) (P16-INK4) (P16-INK4A) (MULTIPLE TUMOR SUPPRESSOR 1) (MTS1). (CDKN2A) | 5 H |  |
|  | U17075; [L36844] | CYCLIN-DEPENDENT KINASE 4 INHIBITOR B (P14-NK4B) (P15-INK4B) (MULTIPLE TUMOR SUPPRESSOR 2) (MTS2) (CDKN2B). | 51 |  |
|  | U40343; [U20498] | CYCLIN-DEPENDENT KINASE 4 INHIBITOR D (P19-INK4D). | 5 J |  |
|  | X92669; [X87843] | CDK-ACTIVATING KINASE ASSEMBLY FACTOA MAT1 (RING FINGER PROTEIN (MAT1) (MENAGE A TROIS) (CDK7/CYCLIN H ASSEMBLY FACTOR) (P36) (P35 (MNAT1) (MAT1) (CAP35). | 5K |  |
|  | U10564 | WEE1-LIKE PROTEIN KINASE (EC 2.7.1.112) (WEE1HU) | 5 L |  |
|  | 001038 | SERINETHREONINE-PROTEIN KINASE PLK (EC 2.7.1.) (PLK-1) (STPK13) | 5M |  |
| ¢ | U38545 | PHOSPHOLIPASE D1 | 5 N |  |
|  | 063878 | NEDD5 PROTEIN HOMOLOG. | 50 |  |
|  | S72008 | CDCIO PROTEIN HOMOLOG | 68 |  |
|  | $\overline{000001}$ | CDC27HS PROTEIN | 6 C |  |
|  | $\underline{\square} 22005$ | UBIQUITIIN-CONJUGATING ENZYME E2-CDC34 | 6 D |  |
|  | U18291 | CDC16HS. | 6 E |  |
|  | 063131 | CDC37 HOMOLOG. | 6 F |  |
|  | U77949 | CDC6-RELATED PROTEIN | 6 G |  |
|  | $\times 60188$ | EXTRACELLULAR SIGNAL-REGULATED KINASE 1 (EC 2.7.1.) (ERK1) (INSULINSTIMULATED MAP2 KINASE) (MAP KINASE 1) (MAPK 1) (P44-ERK1) (ERT2) (P44MAPK) (MICROTUBULE-ASSOCIATED PROTEIN-2 KINASE). | 6 H |  |
|  | M84489 | EXTRACELLULAR SIGNAL-REGULATED KINASE 2 (EC 2.7.1.-) (ERK2) (MITOGENACTIVATED PROTEIN KINASE 2) (MAP KINASE 2) (MAPK 2) (P42-MAPK) (ERT1). | 1 |  |
|  | $\times 80692$ | EXTRACELLULAR SIGNAL-REGULATED KINASE 3 (EC 2.7.1.-) (ERK3) (MAP KINASE ISOFORM P97) (P97-MAPK). | 6 J |  |
|  | $\times 59727$ | EXTRACELLULAR SIGNAL-REGULATED KINASE 4 (EC 2.7.1.) (ERK4) (MAP KINASE ISOFORM P63) (P63-MAPK). | 6 K |  |
|  | $\underline{2} 2278$ | EXTRACELLULAR SIGNAL-REGULATED KINASE 5 (EC 2.7.1.) (ERK5) (ERK4) (BMK1 KINASE) | 6 L |  |


|  | GenBank \# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: | :---: |
|  | X79483 | EXTRACELLULAR SIGNAL-REGULATED KINASE 6 (EC 2.7.1.) (ERK6) (ERK5) | 6M |  |
|  | L35253; [L35263] | MITOGEN-ACTIVATED PROTEIN KINASE P38 (EC 2.7.1.) (MAP KINASE P38) (CYTOKINE SUPPRESSIVE ANTI-INFLAMMATORY DRUG BINDING PROTEIN) (CSAID BINDING PROTEIN) (CSBP) (MAX-INTERACTING PROTEIN 2) (MAP KINASE MXI2). | 6 N |  |
|  | L26318 | STRESS-ACTIVATED PROTEIN KINASE JNK1 (EC 2.7.1.) (C-JUN N-TERMINAL KINASE 1) (JNK-46) | 60 |  |
|  | L31951 | STRESS-ACTIVATED PROTEIN KINASE JNK2 (EC 2.7.1.-) (C-JUN N-TERMINAL KINASE 2) (JNK-55). | 78 |  |
|  | U34819: [U07620] | STRESS-ACTIVATED PROTEIN KINASE JNK3 (EC 2.7.1.-) (C-JUN N-TERMINAL KINASE 3) (JNK3) (MAP KINASE P49 3F12). | 7 C |  |
|  | U25265 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 5 (EC 2.7.1.) (MAP KINASE KINASE 5) (MAPKK 5) (MAPKJERK KINASE 5). | 7 D |  |
|  | L05624 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 1 (EC 2.7.1.-) (MAP KINASE KINASE 1) (MAPKK 1) (ERK ACTIVATOR KINASE 1) (MAPK/ERK KINASE 1) (MEK1). | 7 E |  |
|  | U39657 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 6 (EC 2.7.1.) (MAP KINASE KINASE 6) (MAPKK 6) (MAPKJERK KINASE 6) (SAPKK3) | 7 F |  |
| ¢ | 078876 | MEK KINASE 3 | 7 G |  |
|  | M15796; [J04718] | PCNA (CYCLIN) | 7 H |  |
|  | 449070 | PIN1 | 71 |  |
|  |  | RETINOBLASTOMA-ASSOCIATED PROTEIN [RETINOBLASTOMA |  |  |
|  | M15400 | SUSCEPTIBILITY | 7 J |  |
|  | X74594 | AB2/P130 | 7 K |  |
|  | $\times 74262$ | RBAP48 | 7 L |  |
|  | S66431 | RBP2 RETIINOBLASTOMA BINDING PROTEIN | 7 M |  |
|  | Ṡ57153; S57160 | ṘB P1(RETINOBLASTOMA-BINDING PROTEIN) | 7 N |  |
|  | $\times 85133$ | RBQ1 RETINOPLASTOMA BINDING PROTEIN | 70 |  |
|  | - $\times 15134$ | RBQ-3 | 88 |  |
|  | M96577 | E2F-1 PRB-BINDING PROTEIN | 8 C |  |
|  | Y10479 | E2F-3 | 80 |  |
|  | U15642 | E2F-5 | 8 E |  |
|  | L23959 | E2F-RELATED TRANSCRIPTION FACTOR (DP-1) | 8 F |  |
|  | U18422 | DP2 (HUMDP 2) , DIMERIZATION PARTNER OF E2F | 8 G |  |
|  | U23435: U31089 | ABL INTERACTOR 2 (ABI-2) + ABL BINDING PROTEIN 3 (ABLBP3) [ARGBPIB] | 8 H |  |
|  | -29511 | GRB2 [GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2] | 81 |  |


| GenBank \# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: |
| U69276 | GRB-IR / GRB10 | BJ |  |
| X03484 | RAF ONCOGENE | 8K |  |
| M95712 | RAF,B- | 8L |  |
| J04111 | TRANSCRIPTION FACTOR AP-1 [C-JUN PROTO ONCOGENE] | 8M |  |
| M29039 | JUN B TRANSACTIVATOR | 8 N |  |
| X56681 | TRANSCRIPTION FACTOR JUN-D | 8 O |  |
| M13228 | N-MYC | 9B |  |
| D89667 | C-MYC BINDING PROTEIN | 9 C |  |
| L16785 | NUCLEOSIDE DIPHOSPHATE KINASE B [C-MYC TRANSCRIPTION FACTOR (PUF)] | 9D |  |
| X16416 [M14752] | c-abl | 9 E |  |
|  |  |  |  |
|  | P53 PATHWAY |  |  |
| M14694 | CELLULAR TUMOR ANTIGEN P53 | 9F |  |
| Z12020 | MDM2 PROTEIN (P53-ASSOCIATED PROTEIN) + MDM2-A (GB: U33199) + MDM2-C (GB: U33201) | 9G |  |
| AF007111 | MDM2-LIKE P53-BINDING PROTEIN (MDMX) | 9 H |  |
| Y11416 | P73, A MONOALLELICALLY EXPRESSED P53-RELATED PROTEIN | 91 |  |
| AF010310 AF010311 | P53 INDUCED PROTEIN | 9 J |  |
| AF010309 | PIG3 (PIG3) | 9 K |  |
| AF010312 | PIG7 (PIG7) | 9 L |  |
| AF010314 | PIG10 (PIG10) | 9M |  |
| AF010315 | PIG11 (PIG11) | 9 N |  |
| AF010316 | PIG12 (PIG12) | 90 |  |
| U90313 | GLUTATHIONE-S-TRANSFERASE HOMOLOG | 10B |  |
| U66469 | P53-DEPENDENT CELL GROWTH REGULATOR CGR19 | 10C |  |
| AF001954 | GROWTH INHIBITOR P33ING1 (ING1) | 10D |  |
| L13698 | GROWTH-ARREST-SPECIFIC PROTEIN 1 (GAS-1). . | 10 E |  |
|  |  |  |  |
|  | BCL FAMILY |  |  |
| M14745 | BCL2 | 10F |  |
| U58334 | BCL2 AND P53 BINDING PROTEIN BBP/53BP2 (BBP/53BP2) | 10G |  |
| L22474 | BAX | 10 H |  |
| U59747 | APOPTOSIS REGULATOR BCL-W | 101 |  |
| L08246 | INDUCED MYELOID LEUKEMIA CELL DIFFERENTIATION PROTEIN MCL-1 (ORF IS AT NT. 61-1053; ML) | 10 J |  |


| GenBank \# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: |
| U29680 | BCL2-RELATED PROTEIN A1 (BFL-1 PROTEIN) (HEMOPOIETIC-SPECIFIC EARLY RESPONSE PROTEIN) (GRS PROTEIN) | 10K |  |
| X89986; [U34584] | BCL-2 INTERACTING KILLER (APOPTOSIS INDUCER NBK) (BP4) (BIP1) (BIK) | 10L |  |
| U23765; [U16812; | BCL-2 HOMOLOGOUS ANTAGONIST/KILLER (APOPTOSIS REGULATOR BAK) | 10M |  |
| S82185 | BRAG-1=BRAIN-RELATED APOPTOSIS GENE/BCL-2 HOMOLOG | 10 N |  |
| U66879 | BAD PROTEIN (BCL-2 BINDING COMPONENT 6). | 100 |  |
| S83171; [Z35491] | BCL-2 BINDING ATHANOGENE-1 (BAG-1) (GLUCOCORTICOID RECEPTORASSOCIATED PROTEIN RAP46). | 11B |  |
| U76376 | Harakiri, a protein that activates cell death and interacts w. Bcl-2 and Bcl-XL | 11C |  |
|  | CASPASE CASCADE |  |  |
|  | ..... CASPASES **.. |  |  |
| U13699; (M87507; X6 | (ICE) (INTERLEUKIN-1 BETA CONVERTING ENZYME) (P45) (CASPASE-1) | 110 |  |
| U13021; [U13022] | (CASPASE-2) (ICH-1L) (ICH-1S) | 11E |  |
| U13737 | APOPAIN PRECURSOR (EC 3.4.22.-) (CYSTEINE PROTEASE CPP32) (YAMA PROTEIN) (CASPASE-3) (CPP32) (YAMA PROTEIN) (CASPASE-3) ISOFORM ALPHA | 11F |  |
| U28014; U28015 | ICH-2 PROTEASE PRECURSOR (EC 3.4.22.-) (TX PROTEASE) (ICEREL-II) (CASPASE-4) + CASPASE-5 PRECURSOR (EC 3.4.22.-) (ICH-3 PROTEASE) (TY PROTEASE (ICEREL-III). | 11G |  |
| U20537; U20536 | CASPASE-6 PRECURSOR (EC 3.4.22.-) (APOPTOTIC PROTEASE MCH-2) ISOFORM BETA + ISOFORM ALPHA | 11 H |  |
| U37448 | CASPASE-7 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 3) (ICELAP3) (APOPTOTIC PROTEASE MCH-3) (CMH-1) (LICE2) | 111 |  |
| U60520; U58143; X9\% | CASPASE-8 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 5) (MORT1-ASSOCIATED CED-3 HOMOLOG) (MACH) (FADD-HOMOLOGOUS ICEICED-3-LIKE PROTEASE) (FADD-LIKE ICE) (FLICE) (APOPTOTIC CYSTEINE PROTEASE) (APOPTOTIC PROTEASE MCH-5) (CAP4) (CASP8) (MCH5) ISOF | 11J |  |
| U60520; U58143; X98 | CASPASE-8 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 5) (MORT1-ASSOCIATED CED-3 HOMOLOG) (MACH) (FADD-HOMOLOGOUS ICE/CED-3-LIKE PROTEASE) (FADD-LIKE ICE) (FLICE) (APOPTOTIC CYSTEINE PROTEASE) (APOPTOTIC PROTEASE MCH-5) (CAP4) (CASP8) (MCH5) ISOF | 11K |  |
| U56390; [U60521] | CASPASE-9 PRECURSOR (EC 3.4.22.-) (ICE-LIKE APOPTOTIC PROTEASE 6) (ICELAP6) (APOPTOTIC PROTEASE MCH-6) | 11L |  |
| U60519 | ICE-LIKE APOPTOTIC PROTEASE 4 PRECURSOR (EC 3.4.22.-) (APOPTOTIC PROTEASE MCH-4) (CASPASE-10) | 11M |  |


| GenBank \# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: |
|  | …․ CASPASE REGULATORS ..... |  |  |
| L41690 | TNF RECEPTOR-1 ASSOCIATED PROTEIN (TRADD) | 11 N |  |
| U69108 | TRAF5 | 110 |  |
| U78798; [L81153) | TRAF6 | 128 |  |
| U59863; [U63830] | TRAF-INTERACTING PROTEIN I-TRAF (TRAF FAMILY MEMBER-ASSOCIATED NFKB ACTIVATOR TANK) | 12 C |  |
| U777845 | TRAF-INTERACTING PROTEIN (TRIP) | 12 D |  |
| Y10256 | SERINE THREONINE PROTEIN KINASE, NIK; BINDS SPECIFICALLY TO TRAF2 | 12 E |  |
| AF010127/Y14039; Y | CASPER, A FADD- AND CASPASE-RELATED INDUCER OF APOPTOSIS [CASHALPHA+ CASH-BETA] (FLAME-1) (FLICE-LIKE INHIBITORY PROTEIN) | 12F |  |
| U84388 | DEATH DOMAIN CONTAINING PROTEIN CRADD, APOPTOTIC ADAPTOR MOLECULE FOR CASPASE-2 AND FASLTNF RECEPTOR-INTERACTING PROTEIN RIP | 12G |  |
| U25994; [U50062] | CELL DEATH PROTEIN KINASE RIP | 12 H |  |
| AF015956 | DAXX, A FAS-BINDING PROTEIN THAT ACTIVATES JNK AND APOPTOSIS | 121 |  |
| U12597 | TUMOR NECROSIS FACTOR TYPE 2 RECEPTOR ASSOCIATED PROTEIN (TRAP3) | 12 J |  |
| U21092: [U15637; L3 | CD40 RECEPTOR ASSOCIATED FACTOR 1 (CRAF1) (CAP-1), (LMP1 ASSOCIATED PROTEIN) | 12K |  |
| U45878; [U37546] | INHIBITOR OF APOPTOSIS PROTEIN 1 (HIAP1) (HIAP-1) (C-IAP2) (TNFR2-TRAF SIGNALLING COMPLEX PROTEIN 1) (IAP HOMOLOG C) (IAP1) (MIHC). | 12 L |  |
| U45879: (U37547) | INHIBITOR OF APOPTOSIS PROTEIN 2 (HIAP2) (HIAP-2) (C-IAP1) (TNFR2• TRAF SIGNALLING COMPLEX PROTEIN 2) (IAP HOMOLOG B) (IAP2) (MIHB). | 12M |  |
| U45880; [U32974] | $X$-LINKED INHIBITOR OF APOPTOSIS PROTEIN (X-LINKED IAP) (IAP. LIKEPROTEIN) (HILP). | 12N |  |
|  | LIGANDS AND RECEPTORS |  |  |
| 001394 | TUMOR NECROSIS FACTOR [TNF-a] | 120 |  |
| D12614 | LYMPHOTOXIN-ALPHA [FORMERLY TUMOR NECROSIS FACTOR BETA (TNFb)] | 14B |  |
| L11015 | LYMPHOTOXIN-BETA | 14 C |  |
| U69611 | TNF-ALPHA CONVERTING ENZYME | 140 |  |
| D38122; (U08137) | FAS ANTIGEN LIGAND (APOPTOSIS ANTIGEN LIGAND) (APTL) (APT1LG1) (FASL). | 14 E |  |
| U57059 | APO-2 LIGAND (TNF-RELATED APOPTOSIS INDUCING LIGAND TRAIL) | 14F |  |


|  | GenBank \# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: | :---: |
|  | AF017986 | SECRETED APOPTOSIS RELATED PROTEIN 1 | 14 G |  |
|  | ĀF017988 | SECRETED APOPTOSIS RELATED PROTEIN 3 (SARP3) | 14 H |  |
|  | M33294 | TUMOR NECROSIS FACTOR RECEPTOR TTUMOR NECROSIS FACTOR RECEPTOR 1 (55KD)] | 158 |  |
|  | МЗз294 | TUMOR NECROSIS FACTOR RECEPTOR [TUMOR NECROSIS FACTOR |  |  |
|  | M32315 | RECEPTOR 2] | 15 C |  |
|  | 270519 | FAS/APO 1 | 15 D |  |
|  | U90875 | CYTOTOXIC LIGAND TRAIL RECEPTOR | 15 E |  |
|  | AF016268 | DEATH RECEPTOR 5 (DR5) | 15 F |  |
|  | Y09392:[U75380;U7 | WSL-LR, WSL-S1, WSL-S2 + TRAMP (APO-3) (DDR3) | 15G |  |
|  | M27544 | INSULIN-LIKE GROWTH FACTOR IA | 15 H |  |
|  | M29645 | INSULIN-LIKE GROWTH FACTORII (Somatomedin A] | 168 |  |
|  | $\overline{0} 04434$ | INSULIN-LIKE GROWTH FACTORI RECEPTOR | 16 C |  |
|  | Y00285 [J03528] | CATION-INDEPENDENT MANNOSE-6-PHOSPHATE RECEPTOR linsuline-like rowth fato receptor II IGFR-2 | 16D |  |
|  | D25216 | IGFBP COMPLEX ACID LABILE CHAIN | 16 E |  |
|  | M35410 | IGFBP2 | 16 F |  |
| $\stackrel{\infty}{+}$ |  | IGFBP3 (GROWTH HORMONE-DEPENDENT INSULIN-LIKE GROWTH FACTOR- |  |  |
|  | M31159; [M35878] | BINDING PROTEIN) | 16 G |  |
|  | M65062 | IGFBP5 | 178 |  |
|  | M62402 | IGFBP6 | 17 C |  |
|  |  |  |  |  |
|  |  | OTHER REGULATORS |  |  |
|  | U18321: [ $\times 83544$ ] | DEATH-ASSOCIATED PROTEIN 3 (DAP-3) (ionizing radiation resistance conferring protein) | 17D |  |
|  | X76104 | DEATH-ASSOCIATED PROTEIN KINASE 1 (EC 2.7.1-) (DAP KINASE 1). | 17 E |  |
|  | $\times 86779$ | Fas-activated serine/threonine kinase (FAST) phosphorylates TIA-1 | 17 F |  |
|  | S78085 | PDCD2 | 17G |  |
|  | M63167 | Akt1 (rac protein kinase alpha, protein kinase B, C-Akt) | 17 H |  |
|  | M77198: [M95936] | AKT2 (rac protein kinase beta) | 188 |  |
|  | U63295 | seven in absentia homolog | 18 C |  |
|  | U37688 | RATS1 | 18 D |  |
|  | U91985 | DNA fragmentation factor-45 | 18 E |  |
|  | AFFO23885 | apoptosis-related protein TFAR15 (TFAR15) | 18 F |  |
|  | U56976 | calmodulin dependent phosphodiesterase PDE1B1 | 18G |  |



| GenBank \# | Cell Cycle - Gene Name | Array Coordinate |  |
| :---: | :---: | :---: | :---: |
| D21090 | UV EXCISION REPAIR PROTEIN PROTEIN RAD23 [xeroderma pigmentosum group C repair complementing protein $\mathrm{P} 58 / \mathrm{HHR23B}$ ] | 23G |  |
|  | HOUSEKEEPING GENES |  |  |
| M26880 | UBIQUITIN | 1A |  |
| M86400 | PHOSPHOLIPASE A2 | 1 B |  |
| V00530 | HYPOXANTHINE-GUANINE PHOSPHORIBOSYLTRANSFERASE | 1 C |  |
| X01677 | GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE | 10 |  |
| K00558 | TUBULIN ALPHA | 1 E |  |
| M11886 | HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, C-4 ALPHA CHAIN(MHC) | 1 F |  |
| X00351 | BETA-ACTIN | 1G |  |
| X56932 | 23 kD HIGHLY BASIC PROTEIN | 1 H |  |
| U14971 | RIBOSOMAL PROTEIN S9 | 11 |  |
|  |  |  |  |
|  | NEGATIVE CONTROLS |  |  |
|  | M13 mp18(+) STRAND DNA | 1 J |  |
|  | I-DNA | 1 K |  |
|  | pUC 18 | 1 L |  |
|  |  |  |  |
|  | CALIBRATION MARKERS | 1M1N1O1P |  |
|  |  |  |  |
|  | ORIENTATION MARKERS |  |  |
|  | Dark spots | 2D2G2J2M3A3P6A | 6P9A9P12A12 |
|  | Faint spots | 2A2B2C2E2F2H212 | K2L2N2O2P4 |
|  | Column 13 is blank |  |  |

## Human Stress Array

In the human stress array according to the subject invention. all of the unique polynucleotide probe compositions correspond to genes that are associated with stress responses of human cells, e.g. stress response regulators and effectors. In a specific human stress array of interest, the spots are as provided in Table 5.

| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| K00650 | C-fos |
| M31630 | CAMP RESPONSE ELEMENT BINDING PROTEIN CRE-BPI (CAMP responsive element binding protein 1) |
| M34356 | CREB (ACTIVE TRANSCRIPTION FACTOR) |
| $\times 60188$ | EXTRACELLULAR SIGNAL-REGULATED KINASE 1 (EC 2.7.1.-) (ERKI) (INSULIN- STIMULATED MAP2 KINASE) (MAP KINASE 1) (MAPK 1) (P44-ERKI) (ERI2) (P44-MAPK) (MICROTUBULEASSOCIATED PROTEIN-2 KINASE). |
| M84489 | EXTRACELIULAR SIGNAL-REGULATED KINASE 2 (EC 2.7.1.-) (ERK2) (MITOGEN- ACTIVATED PROTEIN KINASE 2) (MAP KINASE 2) (MAPK 2) (P42-MAPK) (ERII). |
| $\times 80692$ | EXTRACELLULAR SIGNAL-REGULATED KINASE 3 (EC 2.7.1.-) (ERK3) (MAP KINASE ISOFORM P97) (P97-MAPK). |
| X59727; 338873 | EXTRACELLULAR SIGNAL-REGULATED KINASE 4 (EC 2.7.1.-) (ERK4) (MAP KINASE ISOFORM P63) (P63-MAPK). |
| U25278 | EXIRACELLULAR SIGNAL-REGULATED KINASE 5 (EC 2.7.1.-) (ERK5) (ERK4) (BMKI KINASE). |
| X79483 | EXIRACELLULAR SIGNAL-REGULATED KINASE 6 (EC 2.7.1.-) (ERK6) (ERK5). |
| U53442 | MITOGEN-ACTIVATED PROTEIN KINASE P38 BETA (EC 2.7.1.-) (MAP KINASE P38 BETA). |
| L26318 | STRESS-ACTIVATED PROTEIN KINASE JNKI (EC 2.7.1.-) (C-JUN N-TERMINAL KINASE I) (JNK 46) |
| L31951 | STRESS-ACTIVATED PROTEIN KINASE JNK2 (EC 2.7.1.-) (C-JUN N-TERMINAL KINASE 2) (JNK 55). |
| $\begin{aligned} & \text { U25265: (U71087: } \\ & \text { U7 1088) } \end{aligned}$ | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 5 (EC 2.7.1.-)(MAP KINASE KINASE 5) (MAPKK 5) (MAPK/ERK KINASE 5) (MEK5) |
|  | MAP KINASE KINASE MEK5B. |
|  | MAP KINASE KINASE MEK5C |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| L05624 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 1 (EC 2.7.I.-)(MAP KINASE KINASE 1) (MAPKK 1) (ERK ACTIVATOR KINASE 1) (MAPK/ERK KINASE) (MEKI). |
| L11285 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 2 (EC 2.7.1.-)(MAP KINASE KINASE 2) (MAPKK 2) (ERK ACTIVAIOR KINASE 2) (MAPK/ERK KINASE) (MEK2). |
| 439657 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 6 (EC 2.7.1.-)(MAP KINASE KINASE 6) (MAPKK 6) (MAPK/ERK KINASE 6) (SAPKK3). |
| U78876 | MEK KINASE 3 |
| 063780 | STE2O-LIKE KINASE OXIDANT STRESS KINASE (YSKI. STE20 and SPSI RELAIED KINASE) |
| 477129 | SPSI/STE2O HOMOLOGUE, KHS, ACTIVAIOR OFJUN N-TERMINAL KINASE (HSU77129) |
| 007349 | B LYMPHOCYTE GERMINAL CENTER KINASE (HSUO7349) |
| U66464 | HEMATOPOIETIC PROGENITOR KINASE ACIIVAIOR OF SAPK/JNK (HPKI) (HSU66464) |
| AB005216 | NCK ASH AND PHOSHPHOUPASE C GAMMA-BINDING PROTIEN NAP4(AB005216) |
| $\times 17576$ | NCK MELANOMA CYTOPLASMIC SRC HOMOLOGUE (HSNCK) |
| U24153 | SERINE/THREONINE-PROTEIN KINASE PAK-GAMMA (EC 2.7.1.-) (GAMMA-PAK) (P21ACIIVATED KINASE 3) (PAK65) ( $86 / \mathrm{H} 4$ KINASE) (PAK2) PAK3. |
| M35543 | G25K GTP-BINDING PROTEIN, BRAIN ISOFORM (GP) (CDC42 HOMOLOG) CDC42. |
| U12595 | IUMOR NECROSIS FACIOR TYPE I RECEPTOR ASSOCIATED PROTEIN(IRAPI)(HSUI2595) |
| U12596 | IUMOR NECROSIS FACIOR TYPE I RECEPIOR ASSOCIATED PROTEIN(TRAP2) (HSUl2596) |
| X17620 | NUCLEOSIDE DIPHOSPHATE KINASE A (EC 2.7.4.6) (NDK A) (NDP KINASE A) (TUMOR METASTATIC PROCESS-ASSOCIATED PROTEIN) (METASTASIS INHIBITION FACTOR NM23) (NM23-HI). |
| M64673 | $\qquad$ |
| M65217 | HEAT SHOCK FACTOR PROTEIN 2 (HSF 2) (HEAT SHOCK TRANSCRIPTION FACTOR 2)(HSTF 2). |
| 087073 | HEAI SHOCK IRANSCRIPTION FACIOR 4. |
| 134075 | FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (HUMFRAPX) |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| M35663: (U50648) | INTERFERON-INDUCIBLE RNA-DEPENDENT PROTEIN KINASE (P68 KINASE) |
| 007550 | $\qquad$ |
| 086956 | HEAT-SHOCK PROIEIN 110 KD (KIAAO201) |
| $\begin{aligned} & \text { X54079: } \\ & \text { X03900. L39370: } \\ & \text { X16477: Z23090: } \\ & \text { S7457I) } \end{aligned}$ | HEAT SHOCK 27 KD PROTEIN (HSP 27)(STRESS-RESPONSIVE PROTEIN 27)(SRP27)(ESTROGEN-REGULATED 24 KD PROTEIN) ( 28 KD HEAT SHOCK PROTEIN). |
| X61598: 083174 | 47 KD HEAT SHOCK PROTEIN PRECURSOR (COLLAGEN-BINDING PROIEIN I) (COLLIGIN 1) |
|  | Collagen binding protein 2 (HUMCBP2). |
| M11717: (M59828) | HEAT SHOCK 70 KD PROTEIN 1 (HSP70.1) (HSP70-1/HSP70-2). |
| L26336 | HEAT SHOCK-RELATED 70 KD PROTEIN 2 (HEAT SHOCK 70 KD PROTEIN 2). |
| L12723 | HEAT SHOCK 70 KD PROIEIN 4 (HSP7ORY). |
| X51757: M11236 | HEAT SHOCK 70 KD PROTEIN 6 (HEAT SHOCK 70 KD PROTEIN 8'). |
|  | HEAT SHOCK 70 KD PROTEIN 7 (HEAT SHOCK 70 KD PROTEIN 8) (FRAGMEND). |
| Y00371 | HEAI SHOCK COGNATE 71 KD PROTEIN. |
| X07270: (X15183: M27024: M30626: M30627) | HEAT SHOCK PROTEIN HSP 90-ALPHA (HSP 86). |
| M16060 | HEAT SHOCK PROTEIN HSP 90-BETA (HSP 84) (HSP 90) |
| U15590 | HEAT SHOCK PROTEIN 27 (heart) |
| 567070 | HEAT SHOCK PROIEIN HSP 72 HOMOLOG (FRAGMEND). |
| U40992 | HEAT SHOCK PROTEIN HSP4OHEAT SHOCK PROTEIN HSP40 HOMOLOG. |
| L15189 | REGULATED PROTEIN) (GRP 75) (PEPTIDE-BINDING PROTEIN 74) (PBP74) (MORTALIN) (MOT). |
| U28918 | HSC7O-INTERACTING PROTEIN (PROGESTERONE RECEPIOR-ASSOCIATED P48 PROTEIN) |
| D13388 | DNAJ PROTEIN HOMOLOG 2 (DNAJ2 OR HDJ2) |
| $\begin{aligned} & \text { D49547: (D17749: } \\ & \text { D85429) } \end{aligned}$ | HEAT SHOCK PROTEIN 40 |
| M19645 | 78 KD GLUCOSE REGULATED PROTEIN PRECURSOR (GRP 78) (IMMUNOGLOBULIN HEAWY CHAIN BINDING PROTEIN) (BIP) |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| L̄10284: (LI8887: M94859: M98452) | CALNEXIN PRECURSOR (MAJOR HISIOCOMPATIBILITY COMPLEX CLASS I ANTIGENBINDING PROTEIN P88) (P90) (IP90) |
| M84739 | CALRETICULIN PRECURSOR (CRP55) (CALREGULIN) (HACBP) (ERP60)(52 KD RIBONUCLEOPROTEIN AUTOANTIGEN RO/SS-A) |
| j05016 | PROTEIN DISULFIDE ISOMERASE-RELATED PROTEIN PRECURSOR (ERP72) |
| L24804: (L24805) | P23 PROGESTERONE RECEPIOR ASSOCIATED PROTEIN (HUMPRA) |
| M86752 | IRANSFORMATION -SENSITIVE PROTEIN (IEF SSP 3521) |
| 111667 | CYCLOPHILIN-4O |
| U73704 | $48 \mathrm{kDa} \mathrm{FKBP-ASSOCIATED} \mathrm{PROTEIN} \mathrm{FAP48}$ |
| U42031 | 54 KDA PROGESTERONE RECEPIOR-ASSOCIATED PROTEIN FKBP54 |
| M34539: (M80199: M80706:M92423: J05340: X55741: X52220) | FK506-BINDING PROTEIN (FKBP) (FKBPI2) (PEPTIDYL-PROLYL CIS-TRANS ISOMERASE) (PPIASE) (ROTAMASE) |
| M88279 | IMMUNOPHILLIN (FKBPS2) |
| M65128 | RAPAMYCIN-BINDING PROTEIN (FKBP-13) |
| $\begin{aligned} & \times 56134 \text { (M14144: } \\ & 219554 \text { ) } \end{aligned}$ | VIMENTIN. INTERMEDIATE FILAMENT PROTEIN |
| M34664: (M22382) | MIIOCHONDRIAL MATRIX PROTEIN PI PRECURSOR (P6O LYMPHOCYIE PROTEIN) (HSPDI OR HSP60) (CHAPERONIN HOMOLOG) (HUCHA60) (HEAT SHOCK PROTEIN 60) |
| S83171: (235491) | BCL-2 BINDING AIHANOGENE-1 (BAG-1) (GLUCOCORIICOID RECEPIOR-ASSOCIATED PROTEIN RAP46). |
| D23662 | UBIQUITIN-LIKE PROTEIN (NEDD8) |
| X52882 | T-COMPLEX PROTEIN I. ALPHA SUBUNIT (TCP-1-ALPHA)(CCT-ALPHA) CCII OR CCTA OR ICPI |
| U38846 | T-COMPLEX PROTEIN I, DELTA SUBUNIT (TCP-I-DELTA)(CCT-DELTA) (STIMULATOR OF TAR RNA BINDING) (HSU38846). |
| 043950 | T-COMPLEX PROIEIN 1. EPSILON SUBUNIT (ICP-1-EPSILON)(CCT-EPSILON) (HUMKGIDD) |
| x 74801 : (U17104) | I-COMPLEX PROTEIN I. GAMMA SUBUNIT (TCP-I-GAMMA)(CCT-GAMMA) (CCT3) OR (CCIG) OR (IRIC5) (HSHUMAPC). |


| GenBank \# | STRESS RESPONSE REGULAIORS AND EFFECTORS |
| :---: | :---: |
| U83843 | T-COMPLEX PROTEIN I, ETA SUBUNIT (ICP-I-ETA) (CCT-ETA)(HIV-I NEF INTERACTING PROTEIN) (HSU83843). |
| 013627 | T-COMPLEX PROTEIN I, THETA SUBUNIT (TCP-1-THETA)(CCT-THETA) (HUMRSC548). |
| х06985 | HEME OXYGENASE 1 (EC 1.14.99.3) (HO-1) (HSOXYGR). |
| D21243: $(534389)$ | HEME OXYGENASE 2 (EC 1.14.99.3) (HO-2) |
| X15187: (M33716) | ENDOPLASMIN PRECURSOR (94 KD GLUCOSE-REGULATED PROTEIN)(GRP94) (GP96 HOMOLOG) (TUMOR REJECTION ANTIGEN I) (HSTRAI). |
| 005569 | ALPHA CRYSTALLIN A CHAIN (HSUO5569). |
| 545630 | ALPHA CRYSTALLIN B CHAIN (ALPHA (B)-CRYSTALLIN) (ROSENTHAL FIBER COMPONENI). |
| U59058 | BEIA CRYSIALLIN A3 (HSU59058). |
| U59057 | BEIA CRYSTALLIN A4 (HSU59057). |
| U35340 | BEIA CRYSTALLIN BI (CRYBBI) (HSU35340). |
| L10035 | BETA CRYSTALLIN B2 (BP) (HUMCRYB2B). |
| 071216 | BEIA CRYSTALLIN B3 (9CRYBB3 OR CRYB3) (HSU71216). |
| L36869 | BETA CRYSTALLIN S (GAMMA CRYSTALLIN S) (CRYGS) OR (GRYG8). |
| $\begin{aligned} & \text { U66582: M11971: } \\ & \text { (M11970) } \end{aligned}$ | GAMMA CRYSTALLIN C (GAMMA CRYSIALLIN 2 OR I/3) (CRYGC) OR (CRYG3). |
|  | GAMMA CRYSTALLIN B (GAMMA CRYSTALLIN 1-2) (CRYGB) OR (CRYG2) (HUMCRYGXI). |
| L02950 | MU-CRYSTALLIN HOMOLOG (CRYM) (HUMMUCRYS). |
| L13278: (S58039) | QUINONE OXIDOREDUCTASE (EC 1.6.5.5) (NADPH:QUINONE REDUCTASE) (ZETACRYSTALLIN). |
| $\begin{aligned} & \text { D16234: } 249835 ; \\ & \text { D83485: U42068) } \end{aligned}$ | PROBABLE PROIEIN DISULFIDE ISOMERASE ER-60 PRECURSOR (EC 5.3.4.1) (ERP60) (58KDA MICROSOMAL PROTEIN) (phospholipase C-alpha) |
| D49489 | PROTEIN DISULFIDE ISOMERASE P5 PRECURSOR (EC 5.3.4.1) (HUMP5). |
| M75715 | EUKARYOTIC PEPTIDE CHAIN RELEASE FACTOR SUBUNIT I (ERFI) (TB3-I) (CII PROTEIN) RFI. |
| D49490 | PROTEIN DISULFIDE ISOMERASE-RELATED PROTEIN PRECURSOR (EC 5.3.4.1) (PDIR) (HUMPDIR). |
| $\begin{aligned} & \text { j02783: } \\ & \text { (X05130:X07077) } \end{aligned}$ | PROTEIN DISULFIDE ISOMERASE PRECURSOR (PDI) (EC 5.3.4.1) /PROLYL 4-HYDROXYLASE BETA SUBUNIT (EC 1.14.11.2) / CELLULAR THYROID HORMONE BINDING PROTEIN (P55)(HSPRO4HY). |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
|  | Glufathione-insulin transhydrogenase (EC 5.3.4.1 /1.8.4.2): protein-disulfide reductase (glutathlone) (HSGITR). |
| M86737 | SIRUCTURE-SPECIFIC RECOGNITION PROTEIN I (SSRPI) (RECOMBINATION SIGNAL SEQUENCE RECOGNITION PROTEIN) (I160) SSRPI. |
| $\begin{aligned} & \text { X63368: (S37374: } \\ & \text { S37375) } \end{aligned}$ | DNAJ PROTEIN HOMOLOGS HSJIA protein: HSIIB proteln.(HSJ-1)(HSHSJIMR) |
| U65785 | 150 KDA OXYGEN-REGULATED PROTEIN ORPI 50 (HSU65785) |
|  | DNA DAMAGE RESPONSE/REPAIR/RECOMBINAIION |
| $\begin{aligned} & \text { X90392 : (140817: } \\ & \text { U06846) } \end{aligned}$ | MUSCLE-SPECIFIC DNASE I-LIKE (DNase X) (XIB) |
| L24564 | RAD |
| M96684 | IRANSCRIPTIONAL ACTIVAIOR PROTEIN PUR-ALPHA |
| M29971 | METHYLATED-DNA--PROTEIN-CYSTEINE METHYLTRANSFERASE (6-O-METHYLGUANINE-DNA METHYLIRANSFERASE) (MGMD |
| U09579: (L25610) | CYCLIN-DEPENDENT KINASE INHIBITOR I (MELANOMA DIFFERENTIATION ASSOCIATED PROTEIN 6) (MDA-6) (P2I) (CDK-INTERACTING PROTEIN I) (CIPI) (WAFI) (CDKNIA) (CDKNI) (SDII) (PICI) (CAP20) |
| 137374 | FLAP ENDONUCLEASE-1 (MATURATION FACIOR 1) (MFI) (FEN-1) |
| 070310 | DNA REPAIR PROTEIN XRCC9 |
| $\begin{aligned} & \mathrm{HT3218} \text { (X02317: } \\ & \text { K00065) } \\ & \hline \end{aligned}$ | SUPEROXIDE DISMUTASE (CU-ZN) (EC 1.15.1.1) SODI. |
| J02947 | EXIRACELLULAR SUPEROXIDE DISMUTASE PRECURSOR (CU-ZN) (EC 1.15.1.I) (EC-SOD) SOD3. |
| X07834: ( $\times 59445$ ) | SUPEROXIDE DISMUTASE PRECURSOR (MN) (EC 1.15.1.1) SOD2 |
| M14694: (M14695) | CELLULAR IUMOR ANTIGEN P53 |
| 212020: (M92424) | MDM 2 PROTEIN (P53-ASSOCIATED PROTEIN) |
|  | MDM2-A (GB: U33199) |
|  | MDM2-C (GB: U33201) |
| U33841 | ATAXIA TELANGIECTASIA (ATM) |
| J03250 | DNA IOPOISOMERASE I (TOPI) |
| j04088 | DNA IOPOISOMERASE II. ALPHA (IOP2A) |
| $\times 68060$ | DNA TOPOISOMERASE II. BETA (IOP2B) |
| U43431 | DNA TOPOISOMERASE III (TOP3) |


|  | GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: | :---: |
|  | S40706 (S62138) | GROWTH ARREST AND DNA-DAMAGE-INDUCIBLE PROTEIN GADDI53 (DNA-DAMAGE INDUCIBLE TRANSCRIPT 3) (DDIT3) (C/EBP-HOMOLOGOUS PROTEIN) (CHOP) |
|  | $\times 04076$ | CATALASE (EC 1.11.1.6) CAT. |
|  | $\times 51420$ | 5.6-DIHYDROXYINDOLE-2-CARBOXYLIC ACID OXIDASE PRECURSOR (DHICA OXIDASE) (TYROSINASE-RELATED PROTEIN I) (IRP-1) (CATALASE B) (GLYCOPROIEIN-75) (GP75) |
|  |  | BASE EXCISION REPAIR |
|  | X15653 | URACIL-DNA GLYCOSYLASE PRECURSOR (UNG1) |
|  | X52486 | URACIL-DNA GLYCOSYLASE 2 (UNG2) |
|  | M74905 | DNA-3 METHYLADENINE GLYCOSYLASE (3-METHYLADENINE DNA GLYCOSYLASE) (ADPG) (3-ALKYLADENINE DNA GLYCOSYLASE) (N-MEIHYLPURINE-DNA GLYCOSIRASE) (MPG) (MAGI) (3MeAG) |
|  | U51166 | G/T MISMATCH-SPECIFIC THYMINE DNA GLYCOSYLASE (TDG) |
|  | Y11838 | 8.OXYGUANINE DNA GLYCOSYLASE HOMOLOG 1 (muIM HOMOLOG) (OGHI) (HOGGI) (FaPyG) |
| ' | U63329 | mutY HOMOLOG (HMYH) |
|  | X59764: (X66133) | DNA-(APURINIC OR APYRIMIDINIC SITE) LYASE (AP ENDONUCLEASE I) (APEX NUCLEASE) (APEN) (REF-I PROTEIN) (APEI) |
|  | U79718 | ENDONUCLEASE III HOMOLOG 1 (HNTHI) (OCIS3) |
|  | M36067 | DNA LIGASE I (POLYDEOXYRIBONUCLEOTIDE SYNTHASE (ATP)) (DNLI) (LIGI) |
|  | $\times 84740$ | DNA LIGASE III (POLYDEOXYRIBONUCLEOTIDE SYNTHASE (ATP)) (DNL3) |
|  | M18112 | POLY (ADP-RIBOSE) POLYMERASE (PARP) (ADPRI) (NAD (+) ADP-RIBOSYLIRANSFERASE) (POLY (ADP-RIBOSE) SYNTHETASE) (PPOL) |
|  | D16581 | 7.8-DIHYDRO-8-OXOGUANINE TRIPHOSPHATASE (mutT HOMOMOLOG) (8-OXODGIPASE) (MTHI) |
|  | M36089 | DNA-REPAIR PROIEIN XRCCI |
|  | D29013 | DNA POLYMERASE BETA (DPOB) |
|  | M11722 | DNA NUCLEOTIDYLEXOTRANSFERASE (IERMINAL ADDITION ENZYME) (TERMINAL DEOXYNUCLEOTIDYLIRANSFERASE) (TERMINAL TRANSFERASE) (DNT) (IDI) |
|  | $\times 55715$ | 4OS RIBOSOMAL PROIEIN S3 (POSSIBLE dRPOSE) |
|  |  | NUCLEOTIDE EXCISION REPAIR |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| D14533 | DNA-REPAIR PROTEIN COMPLEMENTING XP-A CELLS (XERODERMA PIGMENTOSUM GROUP A COMPLEMENTING PROTEIN) |
| M31899 | DNA-REPAIR PROTEIN COMPLEMENTING XP-B CELLS (XERODERMA PIGMENTOSUM GROUP B COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROTEIN ERCC3) (BASAL IRANSCRIPTION FACIOR 289 KD SUBUNII) (BTF2-D89) (FFIH 89 KD SUBUNII) |
| D21089 | DNA-REPAIR PROTEIN COMPLEMENTING XP-C CELLS (XERODERMA PIGMENTOSUM GROUP C COMPLEMENTING PROTEIN) (pl25) |
| D21235 | UV EXCISION REPAIR PROTEIN PROTEIN RAD23 HOMOLOG A (HHR23A) |
| D21090 | UV EXCISION REPAIR PROTEIN PROTEIN RAD23 HOMOLOG B (HHR23B) (XP-C REPAIR COMPLEMENTING COMPLEX 58 KD PROTEIN) (p58) |
| X52221: (HIII75) | DNA-REPAIR PROTEIN COMPLEMENTING XP-D CELLS (XERODERMA PIGMENTOSUM GROUP D COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROTEIN ERCC-2) |
| U18299 | DAMAGE-SPECIFIC DNA BINDING PROTEIN PI27 SUBUNIT: IMPLICATED IN XERODERMA PIGMENTOSUM GROUP E (DDBI) |
| U18300 | DAMAGE-SPECIFIC DNA BINDING PROTEIN P48 SUBUNIT: IMPLICATED IN XERODERMA PIGMENTOSUM GROUP E (DDB2) |
| $\overline{177890}$ | DNA-REPAIR PROTEIN COMPLEMENTING XP-F CELLS (XERODERMA PIGMENIOSUM GROUP F COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROTEIN ERCC-4) |
| L20046: (X69978) | DNA-REPAIR PROTEIN COMPLEMENTING XP-G CELLS (XERODERMA PIGMENIOSUM GROUP G COMPLEMENTING PROTEIN) (DNA EXCISION REPAIR PROIEIN ERCC-5) |
| U28413 | COCKAYNE SYNDROME GROUP A; WD-REPEAT PROTEIN (CSA PROIEIN) |
| L04791 | EXCISION REPAIR PROTEIN ERCC-6 (CSB) |
| M95809 | BASIC IRANSCRIPTION FACTOR 62 KD SUBUNIT (p62) (BIF2p62) |
| 230094 | BASIC IRANSCRIPTION FACIOR 2. 44 KD SUBUNIT (BIF2P44) |
| 230093 | BASIC IRANSCRIPTION FACTOR 2, 34 KD SUBUNIT (BIF2p34) |
| Y07595 | BȦSIC IRANSCRIPIION FACIOR 2.52 KD SUBUNIT (BTF2p52) |
| M13194 | DNA EXCISION REPAIR PROTEIN ERCC-1 |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| M63488 | REPUCATION PROTEIN A 70 KD DNA-BINDING SUBUNIT (RP-A) (RF-A) (REPLICATION FACIOR-A PROTEIN I) (SINGLE STRANDED DNA-BINDING PROTEIN) |
| J05249 | REPLICATION PROTEIN A 32 KD SUBUNIT (RP-A) (RF-A) (REPLICATION FACIOR-A PROTEIN 2) |
| L07493 | REPUCATION PROTEIN A 14 KD SUBUNIT (RP-A) (RF-A) (REPLCATION FACIOR A PROIEIN 3) |
| U24186 | REPLICATION PROTEIN A 30 KD SUBUNIT (RP-A) (RF-A) (REPLICATION FACTOR-A PROTEIN 4) |
| M15796: (J04718) | PROLIFERATING CELL NUCLEAR ANTIGEN (PCNA) (CYCLIN) |
| L07540 | ACIIVAIOR 136 KD SUBUNIT (REPLICATION FACIOR C 36 KD SUBUNID (RFC36) |
| M87339 | ACTIVAIOR I 37 KD SUBUNIT (REPLICATION FACTOR C 37 KD SUBUNII) (RFC37) |
| L07541 | ACTIVAIOR I 38 KD SUBUNIT (REPLICAIION FACIOR C 38 KD SUBUNII) (RFC38) |
| M87338 | ACIIVAIOR I 40 KD SUBUNIT (REPLICATION FACIOR C 40 KD SUBUNII) (RFC40) |
| L14922 | ACTIVAIOR I 140KD SUBUNIT (REPLCATION FACIOR C LARGE SUBUNII) (AI 140 KD SUBUNII) (RF-C 140 KD SUBUNID) (ACTIVATOR I LARGE SUBUNII) (DNA-BINDING PROTEIN PO-GA) |
| $\times 06745$ | DNA POLYMERASE ALPHA |
| M80397 | DNA POLYMERASE DELIA CATALYTIC CHAIN |
| M60974 | GROWTH ARREST AND DNA-DAMAGE-INDUCIBLE PROTEIN GADD-45 (DNA-DAMAGE INDUCIBLE TRANSCRIPT I) (DDITI) (GA45) |
| S40700 (S62138) | GROWTH ARREST AND DNA-DAMAGE-INDUCIBLE PROTEIN GADDI53 (DNA-DAMAGE INDUCIBLE PROTEIN) (CHOP). |
|  | Homologous recombination |
| U63139 | DNA REPAIR PROTEIN RAD50 |
| D13804: (D14134) | DNA REPAIR PROTEIN RAD5I HOMOLOG |
| U12134 | DNA REPAIR PROTEIN RAD52 HONOLOG |
| प09820 | X-LINKED HELICASE II (X-LINKED NUCLEAR PROTEIN) (XNP) (RAD54L) (XH2) |
| $\times 97795$ | DNA REPAIR PROIEIN RAD54 HOMOLOG |
| 014680 | BREAST CANCER TYPE I SUSCEPTIBILITY PROTEIN (BRCAI) |
| U43746 | BREAST CANCER TYPE 2 SUSCEPTIBILITY PROTEIN (BRCA2) |
| D63882 | MEIOIIC RECOMBINATION PROIEIN DMCI/UM15 HOMOLOG |
| $\times 83441$ | DNA LIGASE IV (POLYDEOXYRIBONUCLEOTIDE SYNTHASE (ATP)) (DNL4) |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| M 74524 | HHR6A (YEAST RAD6 HOMOLOG) (UBIQITIN-CONJUGATING ENZYME) (UBCA) |
| M74525 | HHROB (VEAST RAD6 HOMOLOG) (UBIQIIIN-CONJUGATING ENZYME) (UBCB) |
| Y08837 | RAD5I-LIKE PROTEIN (POSSIBLE XRCC2) |
|  | Non-homologous end-rejoinling |
| U40622 | DNA REPAIR PROTEIN XRCC4 |
| M32865: (S38729) | ATP-DEPENDENT DNA HELICASE II. 70 KD SUBUNIT (LUPUS KU AUTOANTIGEN PROTEIN P70) (70 KD SUBUNIT OF KU ANTIGEN) (THYROID-LUPUS AUTO-ANTIGEN) (TLAA) (KU70) (CIC BOX BINDING FACIOR 75 KD SUBUNII) (CTCBF) (CIC75) (XRCC6) |
| M30938 | ATP-DEPENDENT DNA HELICASE II, 86 KD SUBUNIT (LUPUS KU AUTOANTIGEN PROTEIN P86) ( 86 KD SUBUNIT OF KU ANTIGEN) (THYROID-LUPUS AUTOANTIGEN) (TLAA) (CTC BOX BINDING FACIOR 85 KD SUBUNIT) (CTCBF) (CTC85) (NUCLEAR FACTOR IV) (KU80) (XRCC5) |
| U35835: (U47077) | DNA-DEPENDENT PROTEIN KINASE (DNA-PK) |
|  | DNA DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT (DNA-PKCS) (XRCC7) |
| M29474 | V(D) J RECOMBINATION ACTIVATING PROTEIN I (RAGI) (RAG-1) |
| M94633 | V(D)J RECOMBINATION ACTIVATING PROIEIN 2 (RAG2) (RAG-2) |
|  | MISMATCH REPAIR |
| U07418: (U07343) | DNA MISMATCH REPAIR PROTEIN MLHI (mutL HOMOLOG) |
| U04045: (L47583) | DNA MISMAICH REPAIR PROTEIN MSH2 |
| J04810 | DNA MISMAICH REPAIR PROTEIN MSH3 (DIVERGENT UPSIREAM PROTEIN) (MISMAICH REPAIR PROTEIN 1) (MRPI) (DUP) (DUG) |
| U54777 | DNA MISMATCH REPAIR PROTEIN MSH6 (mutS - ALPHA 160 KD SUBUNII) (G/T MISMATCH BINDING PROTEIN) (GTBP) (GTMBP) (P160) |
| U13696 | DNA MISMAICH REPAIR PROTEIN PMS2 (PMSI PROTEIN HOMOLOG 2) |
| U13695 | DNA MISMAICH REPAIR PROTEIN PMSI (PMSI PROTEIN HOMOLOG 1) |
|  | DRUG/XENOBIOTIC METABOLISM |
| $\times 14672 \times 17059$ | ARYLAMINE N-ACETYLTRANSFERASE, POLYMORPHIC (EC 2.3.1.5) (PNAD) + |
|  | ARYLAMINE N-ACETYLIRANSFERASE, MONOMORPHIC (EC 2.3.1.5) (MNAD) |
| 200036 | CYIOCHROME P450 IA2 (EC 1.14.14.1) (P450-P3) (P450-4).515 |
| 200036 | CYIOCHROME P450 IA2 (EC 1.14.14.1) (P450-P3) (P450-4).515 |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| J04449: D00003: J04813: D00408 | CYTOCHROME P450 IIIA4 (EC 1.14.14.1) (NIFEDIPINE OXIDASE) (NF-25) (P450-PCNI) |
|  | CYTOCHROME P450 IIIA3 (EC 1.14.14.1) (GLUCOCORIICOID-INDUCIBLE) (HLP) CYP3A3. |
|  | CYIOCHROME P450 IIIA5 (EC 1.14.14.1) (P450-PCN3) |
|  | CYTOCHROME P450 IIIA7 (EC 1.14.14.1) (P450-HFLA) |
| J02871 | CYTOCHROME P450 IVBI (EC 1.14.14.1) (P450-HP) |
| $\begin{aligned} & \text { M33318: (X13930: } \\ & \text { X13897): M33317 } \end{aligned}$ | CYOOCHROME P450 IIA6 (EC 1.14.14.1) (COUMARIN 7-HYDROXYLASE) (IIA3) (P450(1)) (PHENOBARBITAL-INDUCIBLE) |
| CYOCHROME P450 IIA7 (EC 1.14.14.1)(P450IIA4) | CYOCHROME P450 IIA7 (EC 1.14.14.1) (P450-IIA4) |
| M21940: M15331: (M21939)M61858: (L07093): M61853: M61854 | CYIOCHROME P450 IIC9 (EC 1.14.14.1) (P450 PB-1) (P450 MP-4) (S-MEPHENYTOIN 4HYDROXYLASE) <br> CYIOCHROME P45O II |
| 009178 | DIHYDROPYRIMIDINE DEHYDROGENASE (NADP + ) PRECURSOR (EC 1.3.1.2) (DPD) (DIHYDROURACIL DEHYDROGENASE) (DIHYDROTHYMINE DEHYDROGENASE) DPYD. |
| M64082 | DIMEIHYLANILINE MONOOXYGENASE (N-OXIDE FORMING) I (EC 1.14.13.8) (FETAL HEPAIIC FLAVIN-CONTAINING MONOOXYGENASE 1) (FMO 1) (DIMEIHYLANILINE OXIDASE 1) |
| M83772 | DIMETHYLANILINE MONOOXYGENASE (N-CXIDE FORMING) 3 (EC I.14.13.8) (HEPAIIC FLAVIN-CONTAINING MONOOXYGENASE 3) (FMO 3) (DIMETHYLANILINE OXIDASE 3) (FMO II) |
| 211737 | DIMEIHYLANILINE MONOOXYGENASE (N-OXIDE FORMING) 4 (EC I.14.13.8) (HEPATIC FLAVIN-CONIAINING MONOOXYGENASE 4) (FMO 4) (DIMETHYLANILINE OXIDASE 4) |
| 137080 | DIMEIHYLANILINE MONOOXYGENASE (N-OXIDE FORMING) 5 (EC I.14.13.8) (HEPATIC FLAVIN-CONTAINING MONOOXYGENASE 5) (FMO 5) (DIMETHYLANILINE OXIDASE 5) |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
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| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| X04808 | PORPHOBILINOGEN DEAMINASE (EC 4.3.1.8) (HYDROXYMETHYLBILANE SYNTHASE) (HMBS) (PRE-UROPORPHYRINOGEN SYNTHASE) |
| M14758 | MULTIDRUG RESISTANCE PROTEIN I (P-GLYCOPROTEIN I) |
| M23234 | MULTIDRUG RESISTANCE PROTEIN 3 (P-GLYCOPROTEIN 3) |
| L05628 | MULIIDRUG RESISTANCE-ASSOCIATED PROTEIN I |
| 008021 | NICOTINAMIDE N-MEIHYLTRANSFERASE (EC 2.1.1.1) |
| $\begin{aligned} & \text { U09031: U28170: } \\ & \text { L19956 } \end{aligned}$ | PHENOL-SULFATING PHENOL SULFOTRANSFERASE 1 (EC 2.8.2.1) (P-PSI) (THERMOSTABLE PHENOL SULFOTRANSFERASE) (IS-PSI) (HASTI/HAST2) (SIIA3) STPI OR STP. |
|  | PHENOL-SULFATING PHENOL SULFOTRANSFERASE 2 (EC 2.8.2.1) (P-PSI) (SIIA2) SIP2. |
|  | MONOAMINE-SULFATING PHENOL SULFOTRANSFERASE (EC 2.8.2.1) (SULFOTRANSFERASE, MONOAMINE-PREFERRING) (M-PSI) (THERMOLABILE PHENOL SULFOTRANSFERASE) (ILPSI (PLACENTAL ESTROGEN SULFOTRANSFERASE) (CATECHOLAMINE-SULFATING PHENOL SULFOTRANSFERASE) (HAST3) STM. |
| U08854: X63359:U06641: J05428:Y00317 | UDP-GLUCURONOSYLTRANSFERASE 2BI5 PRECURSOR. MICROSOMAL (EC 2.4.1.17) (UDPGI) (UDPGTH-3) UGT2BI5. |
|  | UDP-GLUCURONOSYLTRANSFERASE 2BIO PRECURSOR, MICROSOMAL (EC 2.4.1.17) (UDPGT) UGT2BIO. |
|  | UDP-GLUCURONOSYLTRANSFERASE 2B8 PRECURSOR, MICROSOMAL (EC 2.4.1.17) (UDPGD) (ESTRIOL SPECIFIC) (HLUG4) (FRAGMEND) UGT2B8. |
|  | UDP-GLUCURONOSYLTRANSFERASE $2 B 7$ PRECURSOR, MICROSOMAL (EC 2.4.1.17) (UDPGI) (3.4-CATECHOL ESTROGEN SPECIFIC) (UDPGTH-2) UGT2B7. |
|  | UOP-GLUCURONOSYLTRANSFERASE $2 B 4$ PRECURSOR. MICROSOMAL (EC 2.4.1.17) (UDPGI) (HYODEOXYCHOLC ACID) (HLUG25) (UDPGTH-I) UGT2B4. |


|  | GenBank * | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: | :---: |
|  | M68840 | AMINE OXIDASE (FLAVIN-CONTAINING) A (EC I.4.3.4) (MONOAMINE OXIDASE) (MAOA) MAOA. |
|  | M69177 | AMINE OXIDASE (FLAVIN-CONTAINING) B (EC 1.4.3.4) (MONOAMINE OXIDASE) (MAOB) MAOB. |
|  | K03191 | CYTOCHROME P450 IAI (EC 1.14.14.1) (P450-PI) (P450 FORM 6) (P450-C) (TCDDINDUCIBLE). |
|  | $\overline{M 29874}$ | CYTOCHROME P450 IIB6 (EC 1.14.14.1) (PHENOBARBITAL-INDUCIBLE) (P450 \\|BI). |
|  | M20403 | CYTOCHROME P450 IID6 (EC 1.14.14.1) (P450-DBI) (DEBRISOQUINE 4-HYDROXYLASE) CYP2D6. |
|  | J02625 | CYTOCHROME P450 IIE (EC 1.14.14.1) (P450-J) (ETHANOL INDUCIBLE) CYP2EI |
|  | J02906 | CYTOCHROME P450 IFI (EC 1.14.14.1) CYP2FI. |
|  | M14565 | CYOOCHROME P450 XIAI, MITOCHONDRIAL PRECURSOR (EC 1.14.15.6) (P450(SCC)) (CHOLESTEROL SIDE-CHAIN CLEAVAGE ENZYME) (CHOLESIEROL DESMOLASE) CYPIIAI. |
| $\frac{1}{8}$ | X55764 | CYTOCHROME P450 XIBI PRECURSOR (P450C11) (STEROID 11-BETA-HYDROXYLASE) (EC 1.14.15.4) CYP11BI OR S 11 BH. |
|  | M12792: (M23280) | CYTOCHROME P450 XXIB (EC 1.14.99.10) (STEROID 21-HYDROXYLASE) (P450-C21B) CYP21B OR CYP2I OR CYP2IA2. |
|  | 107765 | LIVER CARBOXYLESTERASE PRECURSOR (EC 3.1.1.1) (ACYL COENZYME A:CHOLESTEROL ACYLIRANSFERASE) (ACAD (MONOCYTE/MACROPHAGE SERINE ESTERASE) (HMSE) CES2. |
|  | j05459 | GLUTATHIONE S-TRANSFERASE MU 3 (EC 2.5.1.18) (GSTM3-3) (CLASS-MU) GSTM3 OR GST5. |
|  | 013889 | GLUTATHIONE REDUCIASE |
|  | X 15722 | GLUTATHIONE S-TRANSFERASE MICROSOMAL |
|  | J03746 | GLUTATHIONE S-TRANSFERASE M4 (GLUTATHIONE S-TRANSFERASE MU I) |
|  | $\times 08020$ | GLUTATHIONE S-TRANSFERASE P |
|  | $\times 15480$ | GLUTATHIONE S-TRANSFERASE AI-1 (Glutathione S-transferase (GSI) Ha subunit I) |
|  | M14777 | GLUTATHIONE PEROXIDASE |
|  | M21304 | GLUTHATHIONE S-TRANSFERASE (THEIA 1) |
|  | AFO10310 | GLUIATHIONE-S-TRANSFERASE HOMOLOG |
|  | L05779 | SOLUBLE EPOXIDE HYDROLASE (SEH) (EC 3.3.2.3) (EPOXIDE HYDRATASE) (CYTOSOLIC EPOXIDE HYDROLASE) (CEH) EPHX2. |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| $\overline{\text { M } 57899 ~}$ | UDP-GLUCURONOSYLTRANSFERASE 1-I PRECURSOR. MICROS̄OMAL (EC 2.4.1.17) (UDPGD) (UGT-IA) (UGII•I) (UGII-OI) (UGTI.I) (UGTIAI) (BIURUBIN SPECIFIC ISOZYME 1) (UGIIA) (HUG-BRI) UGII OR GNII. |
| 555985 | UDP-GLUCURONOSYLTRANSFERASE 1-2 PRECURSOR, MICROSOMAL (EC 2.4.1.17) (UDPGI) (UGT-1B) (UGII*2) (UGII-O2) (UGII.2) (UGIIA2) (UGIIB) (HLUGP4) UGII OR GNII. |
| M84127 | UDP-GLUCURONOSYLTRANSFERASE 1-3 PRECURSOR, MICROSOMAL (EC 2.4.1.17) (UDPGD) (UGT-IC) (UGII•3) (UGTI-03) (UGTI.3) (UGIIA3) (UGIIC) UGII OR GNTI. |
| M57951 | UDP-GLUCURONOSYLIRANSFERASE 1-4 PRECURSOR. MICROSOMAL (EC 2.4.1.17) (UDPGD) (UGT-ID) (UGII•4) (UGII-O4) (UGTI.4) (UGTIA4) (UGIID) (BILIRUBIN SPECIFIC ISOZYME 2) (HUG-BR2) UGII OR GNII. |
| J04093 | UDP-GLUCURONOSYLTRANSFERASE 1-6 PRECURSOR, MICROSOMAL (EC 2.4.1.17) (UDPGI) (UGT-IF) (UGII*6) (UGII-O6) (UGII.6) (UGIIA6) (UGIIF) (PHENOL SPECIFIC) UGII OR GNII. |
| X71480 | CYTOCHROME P450 IVAll (EC 1.14.14.1) (FRAGMENI) CYP4A-11. |
| x83573 | ARYLSULFATASE E PRECURSOR (EC 3.1.6.) (ASE) ARSE. |
| X92106 | BLEOMYCIN HYDROLASE (EC 3.4.22.-) (BLM HYDROLASE). |
| M65212 | CATECHOL O-MEIHYLTRANSFERASE. MEMBRANE-BOUND FORM (EC 2.1.1.6) (MB-COMI (CONTAINS: CATECHOL O-MEIHYLTRANSFERASE, SOLUBLE FORM (S-COMD) COMI. |
| 228409 | COPROPORPHYRINOGEN III OXIDASE PRECURSOR (EC 1.3.3.3) (COPROPORPHYRINOGENASE) (COPROGEN OXIDASE) (COX) CPO. |
| Y09501 | NADH-CYTOCHROME B5 REDUCTASE (EC 1.6.2.2) (B5R) DIAI. |
| U12778 | ACYL-COA DEHYDROGENASE, SHORT/BRANCHED CHAIN SPECIFIC PRECURSOR (EC 1.3.99.-) (SBCAD) (2-METHYL BRANCHED CHAIN ACYL-COA DEHYDROGENASE) (2MEBCAD) ACADSB. |
| M74542 | ALDEHYDE DEHYDROGENASE, DIMERIC NADP-PREFERRING (EC 1.2.1.5) (CLASS 3) ALDH3. |


| GenBank \# | STRESS RESPONSE REGULATORS AND EFFECTORS |
| :---: | :---: |
| X 53463 | GLUTATHIONE PEROXIDASE-GASTROINTESTINAL (EC I.11.1.9) (GSHPX-GI) (GLUTATHIONE PEROXIDASE-RELATED PROTEIN 2) (GPRP) GPX2. |
| X71973 | PHOSPHOLIPID HYDROPEROXIDE GLUTHATIONE PEROXIDASE (EC 1.11.1.9) (PHGPX) GPX4. |
| M63012 | SERUM PARAOXONASE/ARYLESTERASE I (EC 3.1.1.2) (EC 3.1.8.1) (PON I) (SERUM ARYLDIAKYLPHOSPHATASE I) (A-ESTERASE I) (AROMATIC ESTERASE I) PONI OR PON. |
| L48513 | SERUM PARAOXONASE/ARYLESTERASE 2 (EC 3.1.1.2) (EC 3.1.8.1) (PON 2) (SERUM ARYLDIAKYLPHOSPHATASE 2) (A-ESTERASE 2) (AROMATIC ESTERASE 2) PON2. |
| L48516 | SERUM PARAOXONASE/ARYLESTERASE 3 (EC 3.1.1.2) (EC 3.1.8.1) (PON 3) (SERUM ARYLDIAKYLPHOSPHATASE 3) (A-ESTERASE 3) (AROMATIC ESTERASE 3) (FRAGMEND) PON3. |
| 562904 | IHIOPURINE S-METHYLTRANSFERASE (EC 2.1.1.67) (HHIOPURINE METHYLIRANSFERASE) TPMT. |
| L02932 | PEROXISOME PROLIFERAIOR ACTIVATED RECEPTOR ALPHA (PPAR-ALPHA) PPARA OR PPAR |
| 107592 | PEROXISOME PROLIFERATOR ACIIVATED RECEPTOR BETA (PPAR-BETA) (PPAR-DELTA) (NUCLEAR HORMONE RECEPIOR I) (NUCI) (NUCI) PPARB OR PPARD. |
|  | HOUSEKEEPING GENES |
| M26880 | UBIQUITIN |
| M86400 | PHOSPHOLIPASE A2 |
| V00530 | HYPOXANTHINE-GUANINE PHOSPHORIBOSYLIRANSFERASE |
| $\times 01677$ | GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE |
| K00558 | IUBULIN ALPHA |
| M11886 | HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, C-4 ALPHA CHAIN |
| (MHC) | 902 |
| $\times 00351$ | BEIA-ACIIN |
| X56932 | 23 kD HIGHLY BASIC PROTEIN |
| U14971 | RIBOSOMAL PROTEIN S9 |
|  |  |
|  | NEGATIVE CONTROLS |

Oncogene and Tumor Suppressor Gene Array
In the oncogene and tumor suppressor gene array according to the subject invention, all of the unique polynucleotide probe compositions correspond to genes that are associated with cellular proliferative diseases, specifically neoplastic diseases. Genes of interest that may be represented on the array include: oncogenes and tumor suppressor genes. In a specific oncogene and tumor suppressor gene array of interest, the spots are as provided in Table 6.


| $\frac{1}{O}$ | GenBank\# | Gene Name |
| :---: | :---: | :---: |
|  | L12260 | HEREGULIN ALPHA [Recombinant glial growth factor 2] |
|  | L12261 | HEREGULIN ALPHA [Recombinant glial growth factor] |
|  | M27288 | ONCOSTATIN M |
|  | M59964 | STEM CELL FACTOR (C-KIT LIGAND) |
|  | M76125 | AXL (TYROSINE-PROTEIN KINASE RECEPTOR UFO) |
|  | X06182 | C-KIT PROTO-ONCOGENE [maststem cell growth factor receptor] |
|  | $\times 06374$ | PLATELET-DERIVED GROWTH FACTOR A CHAIN |
|  | D13866 | ALPHA-CATENIN |
|  | D17517 | SKY (DTK) (TYRO3) (RSE) |
|  | L11353: Z22664; X72657; L27133 | MERLIN (SCHWANNOMIN) (moesin-ezrin-radixin-like protein)(neurofibromatosis 2) |
|  | L13738 | TYROSINE-PROTEIN KINASE SYK [activated p21cdc42Hs kinase (ack)] |
|  | L14837 | TIGHT JUNCTION PROTEIN ZO-1 |
|  | L16785 | NUCLEOSIDE DIPHOSPHATE KINASE B [c-myc transcription factor (puf)] |
|  | L19067 | TRANSCRIPTION FACTOR P65 |
|  | L20422 | PROTEIN ETA [14-3-3 PROTEIN ETA] |
|  | L22075 | GUANINE NUCLEOTIDE REGULATORY PROTEIN (G13) |
|  | L25259 | T LYMPHOCYTE ACTIVATION ANTIGEN CD86 [CD28 antigen ligand 2, 87-2 antigen] |
|  | L33264 | CDC2-RELATED KINASE PISSLRE |
|  | M13150 | MAS PROTO-ONCOGENE |
|  | M31213: [M57464] | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE RECEPTOR RET PRECURSOR (EC 2.7.1.112) (C-RET). (Papillary thyroid carcinoma-encoded protein] |
|  | M31899 | DNA-REPAIR PROTEIN COMPLEMENTING XP-B CELLS [DNA repair helicase (ERCC3)] |
|  | M32865 | ATP-DEPENDENT DNA HELICASE II ( 70 KD SUBUNIT) [Thyroid autoantigen 70kD (Ku antigen)] |
|  | M34960 | TRANSCRIPTION FACTOR IID |
|  | M36089 | DNA-REPAIR PROTEIN XRCC1 |
|  | M54915 | PIM-1 PROTO-ONCOGENE (SERINETHREONINE-PROTEIN KINASE) |
|  | M60915 | NEUROFIBROMIN [neurolibromatosis protein type I (NF1)] |
|  | M62397 | COLORECTAL MUTANT CANCER PROTEIN |


|  | GenBank \# | Gene Name |
| :---: | :---: | :---: |
|  | M62810 | MITF1 [TANSCRIPTION FACTOR 1 MITOCHONDRIAL] |
|  | M81750 | MYELOID CELL NUCLEAR DIFFERENTIATION ANTIGEN |
|  | M81840 | TRANSFORMING PROTEIN MAF [NRL gene product] |
|  | M83234 | Y BOX BINDING PROTEIN-1 [Nuclease-sensitive element DNA-binding protein] |
|  | U02082 | GUANINE NUCLEOTIDE REGULATORY PROTEIN TIM1 |
|  | 003056 | HYALURONIDASE [tumor suppressor (LUCA-1)] |
|  | U07236 | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE LCK [Lymphocyte-specific protein tyrosine kinasel |
|  | U09579: [L25610] | CYCLIN-DEPENDENT KINASE INHIBITOR 1 (MELANOMA DIFFERENTIATION ASSOCIATED PROTEIN 6) (MDA-6) (P21) (CDK-INTERACTING PROTEIN 1) (CIP1) (WAF1) (CDKN1A) (CDKN1) (SD11) (PIC1) (CAP20) |
|  | X07024 | TRANSCRIPTION INITIATION FACTOR TFIID ( 250 KD SUBUNIT) [CG1 protein inv. in cell proliferation) |
|  | X15218 | SKI ONCOGENE |
|  | $\times 15219$ | SKI-RELATED ONCOGENE SNON |
|  | X51630 | WILMS TUMOR PROTEIN |
|  | M81933 | cdc25A; M-PHASE INDUCER PHOSPHATASE 1 (EC 3.1.3.48) |
|  | M92287 | CYCLIN D3 |
|  | S85655 | PROHIBITIN |
|  | $\times 03484$ | RAF PROTO-ONCOGENE (SERINETHREONINE-PROTEIN KINASE) |
|  | $\overline{\times 16416}$ | PROTO-ONCOGENE TYROSINE.PROTEIN KINASE ABL |
|  | Х $\overline{\text { 59798; [M64349] }}$ | CYCLIN D1 (CYCLIN PRAD1) (BCL-1 ONCOGENE) |
|  | D13639 [M90813] | CYCLIN D2 |
|  | $\begin{aligned} & \text { HT2291; [K03214; } \\ & \text { X03996] } \end{aligned}$ | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE SRC (EC 2.7.1.112) (P60-SRC) (C- SRC). |
|  | X75042 | C-REL PROTO-ONCOGENE PROTEIN |
|  | L25080 | TRANSFORMING PROTEIN RHOA [proto-oncogene rhoA, multidrug resislance protein] |
|  | X75342 | SHB ADAPTOR PROTEIN [A SrC HOMOLOGY 2 PROTEIN] |
|  | L26584 | CDC25 [GUANINE NUCLEOTIDE RELEASING PROTEIN] |
|  | X76132 | TUMOR SUPPRESSOR PROTEIN DCC |


|  | GenBank \# | Gene Name |
| :---: | :---: | :---: |
|  | L27211 | CYCLIN-DEPENDENT KINASE 4 INHIBITOR A (CDK4I) (P16-INK4) (P16-INK4A) (MULTIPLE TUMOR SUPPRESSOR 1) (MTS1). (CDKN2A) |
|  | M13228 | N-MYC PROTO-ONCOGENE PROTEIN |
|  | M15400 | RETINOBLASTOMA-ASSOCIATED PROTEIN [retinoblastoma susceptibility] |
|  | M15990 | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE YES |
|  | M19720 | L-MYC-2 PROTEIN |
|  | M19722 | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE FGR (EC 2.7.1.112) (P55-FGR) (C- FGR). |
|  | M73812 | CYCLINE (G1/S-SPECIFIC) |
|  | M74088 | ADENOMATOUS POLYPOSIS COLI PROTEIN |
|  | U25994 | TYROSINE-PROTEIN KINASE LYN [cell death protein RIP] |
|  | U40343: [U20498] | CYCLIN-DEPENDENT KINASE 4 INHIBITOR D (P19-INK4D). |
|  | U43746 | BREAST CANCER TYPE 2 SUSCEPTIBILITY PROTEIN |
|  | X02751 | TRANSFORMING PROTEIN P21 [ N -ras] |
|  | X16706 | FRA-2 [fos-related antigen 2] |
|  | X16707 | FRA-1 [fos-related antigen 1] |
|  | X51521 | EZRIN [Villin 2] |
|  | X56681 | TRANSCRIPTION FACTOR JUN-D |
|  | X59932 | TYROSINE-PROTEIN KINASE CSK [C-SRC-kinase] |
|  | X86779 | FAST KINASE |
|  | X87838 | BETA-CATENIN |
|  | 229090 | PHOSPHATIDYLINOSITOL 3-KINASE CATALYTIC SUBUNIT ALPHA ISOFORM |
|  | M14745 | BCL2 |
|  | D38305 | TOB |
|  | L16464 | ETS-RELATED PROTEIN PE-1 [ETS oncogene (PEP1)] |
|  | L29216 | PROTEIN KINASE CLK (CLK2) |
|  | L29220 | PROTEIN KINASE CLK (CLK3) |
|  | L29222 | PROTEIN KINASE CLK (CLK1) |
|  | U10564 | CDK TYROSINE 15-KINASE WEEIHU |



| $\dot{\square}$ | GenBank\# | Gene Name |
| :---: | :---: | :---: |
|  | 443318 | frizzled 5 |
|  | U46461 | dishevelled homolog (DVL) |
|  | U49262:[U75651] | dishevelled (DVL) + dishevelled 3 (DVL3) |
|  | L34075 | FKBP-RAPAMYSIN ASSOCIATED PROTEIN (FRAP) |
|  | $\times 07876$ | WNT2 OR IRP |
|  | L40027 | glycogen synthase kinase 3 |
|  | $\times 66360$ | SERINETTHREONINE-PROTEIN KINASE PCTAIRE-2 |
|  | $\times 66362$ | SERINETHREONINE PROTEIN KINASE PCTAIRE-3 |
|  | $\times 66363$ | SERINETHREONINE-PROTEIN KINASE PCTAIRE-1 |
|  | X74594 | RB2/p130 |
|  | $\times 85134$ | RBQ-3 |
|  | 271621 | Wnt-13 |
|  | AB000220 | semaphorin E |
|  | AF001954 | growth inhibitor P331NG1 (ING1) |
|  | AF007111 | MDM2-like p53-binding protein (MDMX) |
|  | D89667 | C-myc binding protein |
|  | U29343 | HYALURONAN RECEPTOR (RHAMM) |
|  | 066469 | p53-dependent cell growth regulator CGR19 |
|  | 076638 | BRCA1-ASSOCIATED RING DOMAIN PROTEIN |
|  | U82169 | trizzled homolog (FZD3) |
|  | U84401 | smoothened |
|  | 090875 | cytotoxic ligand TRAIL receptor |
|  | 095299 | Notch4 |
|  | $Y 11416$ | p73, a monoallelically expressed p53-related protein |
|  | X91940 | WNT-8B |
|  | $\times 97057$ | WNT-10B |
|  | Y 10479 | E2F-3 |
|  | Y11306 | beta cateniNTCF-4 |
|  | U38276 | SEMAPHORIN-1 |
|  | U77493 | Notch2 |
|  | K00650 | C-10s |
|  | $\overline{\text { x } 53795}$ | CD82 ANTIGEN (INDUCIBLE MEMBRANE PROTEIN R2) (C33 ANTIGEN) (IA4) (METASTASIS SUPPRESSOR KANGAI 1) (SUPPRESSOR OF TUMORIGENICITY-6). |
|  | L38518 | sonic hedgehog (SHH) |
|  | M 54968 | $\bar{K}-\overline{A C S}$, ONCOGENE |


| $\frac{-1}{9}$ | GenBank \# | Gene Name |
| :---: | :---: | :---: |
|  | M63167 | Akt1 (rac protein kinase alpha, protein kinase B, c-Akt) |
|  | S57153; S57160 | RBP1(RETINOBLASTOMA-BINDING PROTEIN) |
|  | U23435; U31089 | Abl interactor 2 (Abi-2) + Abl binding protein 3 (AbIBP3) [ArgBPIB] |
|  | M96577 | E2F-1 pRB-binding protein |
|  | $\begin{aligned} & \text { U24163; [U91903; } \\ & \text { U68057] } \end{aligned}$ | Irizzled-related FrzB (Fritz) (irezzled (fre)) |
|  | L05148 | TYROSINE-PROTEIN KINASE ZAP-70 (EC 2.7.1.112) (70 KD ZETA-ASSOCIATED PROTEIN) (ZAP70) |
|  | M97935 | SIGNAL TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION 1-ALPHABETA (TRANSCRIPTION FACTOR ISGF-3 COMPONENTS P91/P84) (STAT1) |
|  | U10087 $\times 58957$ | TYROSINE-PROTEIN KINASE BTK (EC 2.7.1.112) (BRUTON'S TYROSINE KINASE)(AGAMMAGLOBULINAEMIA TYROSINE KINASE) (ATK) (B CELL PROGENITOR KINASE) (BPK) (BTK) (AGMX1) |
|  | AF0-16268 | death receptor 5 (DR5) |
|  | M35296 | TYROSINE-PROTEIN KINASE ABL2 (EC 2.7.1.112) (TYROSINE KINASE ARG) (ABLL) |
|  | U18671 M97934 | SIGNAL TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION 2 (P113) (STAT2) |
|  | U47686 | SIGNAL TRANSDUCER AND TRANSCRIPTION ACTIVATOR 5B (STAT5B) |
|  | M80629 | CDC2-RELATED PROTEIN KINASE CHED |
|  | S66431 | RBP2 retinoblastoma binding protein |
|  | U04045: [L47583] | DNA MISMATCH REPAIR PROTEIN MSH2 |
|  | U29656 | DR-NM23 |
|  | U43148 | patched homolog (PTC) |
|  | J02958 | MET |
|  | U49089 | neuroendocrine-dlg (NE-dig) a novel human homolog of the Drosophila discs large (dig) tumor suppressor protein interacting with the APC protein |
|  | U54777 | DNA MISMATCH REPAIR PROTEIN MSH6 (mutS - ALPHA 160 KD SUBUNIT) (G/T MISMATCH BINDING PROTEIN) (GTBP) (GTMBP) (P160) |
|  | X66358 | SERINETHREONINE-PROTEIN KINASE KKIALRE |

## Cell-Cell Interaction Array

In the cell-cell interaction array according to the subject invention, all of the unique polynucleotide probe compositions correspond to genes that are associated with cell-cell interaction, e.g. cell-cell signaling. In a specific cell-cell interaction array of interest. the spots are as provided in Table 7.

| $\frac{1}{1!}$ | GenBank \# | CELL INTERACTION (Gene Names) |
| :---: | :---: | :---: |
|  | M32315 | TUMOR NECROSIS FACTOR RECEPTOR [Tumor necrosis factor receptor 2] |
|  | $\times 01394$ | TUMOR NECROSISFACTOR [TNFa) |
|  | D12614 | LYMPHOTOXIN-ALPHA [formerly tumor necrosis factor beta (TNF-beta)] |
|  | M12807 | T-CELL SURFACE GLYCOPROTEINCD4 |
|  | M14648 | VITRONECTIN RECEPTOR ALPHA [Integrin, alpha V; antigen CD51] |
|  | $\times 75208$ | TYROSINE-PROTEIN KINASE RECEPTOR EPH-3 |
|  | $\times 74764$ | TYROSINE-PROTEIN KINASE CAK [Tyrosine kinase, receptor TKT] |
|  | M18391 | TYROSINE-PROTEIN KINASE RECEPTOR EPH |
|  | $\begin{aligned} & \mathrm{U} 08839 \text { [M83246; } \\ & \times 51675] \end{aligned}$ | UROKINASE PLASMINOGEN ACTIVATOR SURFACE RECEPTOR, GPI-ANCHORED FORM PRECURSOR (U-PAR) (MONOCYTE ACTIVATION ANTIGEN MO3) (CD87 ANTIGEN) |
|  | M33294 | TUMOR NECROSIS FACTOR RECEPTOR [Tumor necrosis factor receptor 1 ( 55 kD )] |
|  | Y00285 | CATION-INDEPENDENT MANNOSE-6-PHOSPHATE RECEPTOR [insuline-like growth factor receptor II, IGFR-2] |
|  | L07414 | CD40 |
|  | L08096; [S69339] | CD27 (CD70 ANTIGEN) |
|  | L09753 | CD30 |
|  | M35410 | IGFBP-2 [INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 2 |
|  | M63928 | CD27L RECEPTOR [T cell activation antigen (CD27)] |
|  | M67454 | FASL RECEPTOR [Fas antigen, APO-1 antigen] |
|  | M83554 | CD30L RECEPTOR [Lymphocyte activation antigen CD30; Ki-1 antigen ] |
|  | X60592 | CD40L RECEPTOR [Cdw40 nerve growth factor receptor-related B-Iymphocyte activation molecule] |
|  | $\begin{aligned} & \text { D13866 [D14705 } \\ & \text { L23805; L22080] } \end{aligned}$ | ALPHA-CATENIN (CADHERIN-ASSOCIATED PROTEIN) (ALPHA E-CATENIN) |
|  | $\begin{aligned} & \text { D25303; } \\ & \text { L24158 } \end{aligned}$ | integrin alpha9 |
|  | J03132 | INTERCELLULAR ADHESION MOLECULE-1 |
|  | J04536 | LEUKOSIALIN [sialophorin (CD43)] |
|  | $\begin{aligned} & \text { L11353; Z22664; } \\ & \times 72657 ; L 27133 \end{aligned}$ | MERLIN (SCHWANNOMIN) (moesin-e2rin-radixin-like protein)(neurofibromatosis 2) |
|  | L13616 | Focal adhesion kinase |
|  | L14837 | TIGHT JUNCTION PROTEIN ZO-1 |
|  | $\begin{aligned} & \text { L16785; } \\ & \text { IM36981] } \end{aligned}$ | NUCLEOSIDE DIPHOSPHATE KINASE B (EC 2.7.4.6) (NDK B) (NDP KINASE B) (NM23-H2) (C-MYC PURINE-BINDING TRANSCRIPTION FACTOR PUF). |


|  | GenBank \# | CELL INTERACTION (Gene Names) |
| :---: | :---: | :---: |
|  | L20815 | $S$ PROTEIN |
|  | $\underline{25259}$ | T LYMPHOCYTE ACTIVATION ANTIGEN CD86_ [CD28 antigen ligand 2, B7-2 antigen] |
|  | L34774 | opioid binding cell adhesion molecule |
|  | M15476 | UROKINASE-TYPE PLASMINOGEN ACTIVATOR PRECURSOR (EC 3.4.21.73) (UPA) (UPLASMINOGEN ACTIVATOR) |
|  | $\begin{aligned} & \text { M15518; [ } \\ & \text { X07393; M18182] } \end{aligned}$ | TISSUE-TYPE PLASMINOGEN ACTIVATOR PRECURSOR (EC 3.4.21.68) (T-PA) (TPLASMINOGEN ACTIVATOR). |
|  | $\begin{aligned} & \text { M18082;[ } \\ & \text { J02685 } \end{aligned}$ | PLASMINOGEN ACTIVATOR INHIBITOR-2, PLACENTAL (PAI-2) (MONOCYTE ARGSERPIN) (UROKINASE INHIBITOR). |
|  | M21097 | CD19 B-LYMPHOCYTE ANTIGEN [Differentiation antigen (CD19)] |
|  | M23197 | CD33 MYELOID CELL SURFACE ANTIGEN [Differentiation antigen (CD33)] |
|  | M28882 | CELL SURFACE GLYCOPROTEIN MUC18 |
|  | M30257 | VASCULAR CELL ADHESION PROTEIN [vascular cell adhesion molecule 1] |
|  | M30640 | E-SELECTIN [Endothelial leucocyte adhesion molecule I (ELAM1)] |
|  | M34064 [X57548; X54315; S42303] | CADHERIN-2 (N-CADHERIN) |
|  | M54992 | CD72 B-CELL DIFFERENTIATION ANTIGEN |
|  | M59040 | CD44 ANTIGEN HEMATOPOIETIC FORM [Cell adhesion molecule (CD44)] |
|  | M63618 | bullous pemphigoid antigen |
|  | M74387 | LICAM |
|  | M74777 | CD26 JDIPEPTIDYL PEPTIDASE IV; adenosine deaminase complexing protein 2] |
|  | 001160 | SAS (TRANSMEMBRANE 4 SUPERFAMILY PROTEIN) |
|  | 03056 | HYALURONIDASE [ ${ }^{\text {lumor suppressor (LUCA-1)] }}$ |
|  | 007819 | CONTACTIN [Contactin 1 (CNTN1)] |
|  | U15979 | DELTA.LIKE PROTEIN [dlk] |
|  | X16841 | N-CAM INEURAL CELL ADHESION MOLECULE, PHOSPHATIDYLINOSITOL-LINKED ISOFORM; CD56] |
|  | $\times 70326$ | MacMarcks |
|  | $\times 74979$ | TYROSINE-PROTEIN KINASE CAK [EDDR1; TRK E] |
|  | Z26317 [S64273] | desmoglein 2 |
|  | L25080 | TRANSFORMING PROTEIN RHOA [proto-oncogene rhoA, multidrug resistance protein] |
|  | $\times 76132$ | DCC |
|  | J02703 | PLATELET MEMBRANE GLYCOPROTEIN IIIA |


|  | GEenBank \# | CELL INTERACTION (Gene Names) |
| :---: | :---: | :---: |
|  | $J 04145$ | INTEGRIN ALPHA M [Neutrophil adherence receptor alpha-M subunit; Complement component receptor 3, alpha; also known as CD11b (p170), macrophage antigen alpha polypeptide] |
|  | J05633 | integrin beta5 |
|  | $\begin{aligned} & 12002 ; \\ & {[\times 16983]} \end{aligned}$ | integrin alpha4 |
|  | $\underline{L 2851}$ | integrin alphaE |
|  | L36531 | integrin alpha8 |
|  | M15395 | LEUKOCYTE ADHESION PROTEIN [CELL SURFACE ADHESION GLYCOPROTEINS LFA1, CR3 AND P150,95, BETA-SUBUNIT] |
|  | $\begin{aligned} & \text { M28249; } \\ & {[\times 17033]} \end{aligned}$ | integrin alpha2 [very late antigen-2 (vla-2)/collagen receptor alpha-2 subunit] |
|  | M34480 | INTEGRIN ALPHA 2B [PLATELET MEMBRANE GLYCOPROTEIN IIB (GPIIb); antigen CD41B] |
|  | M35198 | integrin beta6 |
|  | M59911 | integrin alpha3 |
|  | M62880 | integrin beta7 |
| 二 | M73780 | integrin beta8 |
| A | M81695 | INTEGRIN ALPHA X (LEUKOCYTE ADHESION GLYCOPROTEIN P150,95 ALPHA CHAIN; antigen CD11C (p150)] |
|  | X06256 | integrin alpha5 [fibronectin receptor alpha subunit] |
|  | $\times 07979$ | FIBRONECTIN RECEPTOR (BETA SUBUNIT) [INTEGRIN BETA 1] |
|  | $\begin{aligned} & \times 53586 ; \\ & \times 595121 \end{aligned}$ | integrin alpha6 |
|  | $\begin{aligned} & \times 53587 ; \\ & \times 52186] \end{aligned}$ | integrin beta4 |
|  | $\times 68742$ | integrin alpha |
|  | X74295 | integrin alpha7B |
|  | Y00796 | INTEGRIN ALPHA L [LEUKOCYTE ADHESION GLYCOPROTEIN LFA-1 ALPHA CHAIN; antigen CD11A (p180)] |
|  | D38122 | FAS ANTIGEN LIGAND |
|  | $\begin{aligned} & \text { M74088; } \\ & \text { [M73548] } \end{aligned}$ | APC (DP2.5) |
|  | $\begin{aligned} & \text { U43522; } \\ & \text { L49207] } \end{aligned}$ | Protein tyrosine kinase Pyk2 (Cell adhesion kinase-beta, CAK-beta) (FAK2) |
|  | $\times 51521$ | Ezrin (cytovillin 2) |


| $\frac{1}{4}$ | \|GenBank \# | CELL INTERACTION (Gene Names) |
| :---: | :---: | :---: |
|  | X87838 [219054] | BETA-CATENIN |
|  | L11015 | LYMPHOTOXIN-BETA |
|  | U57059 | FȦS ANTIGEN LIGAND [TNF-related apoptosis inducing ligand TRAIL; Apo-2 ligand] |
|  | D45132 | ANNEXIN I [zinc finger protein RIZ] |
|  | M68516; | PLAASMA SERINE PROTEASE INHIBITOR PRECURSOR (PCI) (PROTEIN C INHIBITOR) (PLASMINOGEN ACTIVATOR INHIBITOR-3) (PAI3). |
|  | [J02639] | Integrin-linked kinase (ILK) |
|  | U43408 | FOCAL ADHESION KINASE [tyrosine kinase (Tnk1)] |
|  | U60800 | semaphorin (CD100) |
|  | U61262 | TUMOR SUPPRESSOR PROTEIN DCC [neogenin] |
|  | L11370 | protocadherin 42 |
|  | $\times 78817$ | RHO-GAP HEMATOPOIETIC PROTEIN C1 (P115) (KIAA0131). |
|  | $\times 85978$ | TAX1, AXONIN-1/TAQ1 |
|  | L11373 | protocadherin 43 |
|  | X89576 | MMP-17 (MT4-MMP) |
|  | Y00815 | LAR |
|  | 230183 | TIMP-3 (mitogen-inducible gene 5, mig-5) |
|  | 235227 | ras-like small GTPase TTF |
|  | $\begin{aligned} & \hline \text { D26512, } \\ & {[\times 83535]} \end{aligned}$ | MMP-14 (MT1-MMP) |
|  | D31784 | CADHERIN-6 |
|  | D50477 | MMP-16 (MT3-MMP) |
|  | D83542 | CADHERIN-14 MUSCLE-CADHERIN PRECURSOR (M-CADHERIN) (CADHERIN-14) (CADHERIN-15) |
|  | J03210, [J05471] | MMP-2 (gelatinase A) |
|  | J05070, [D10051] | MMP-9 (gelatinase B) |
|  | $J 05556$ | MMP-8 (collagenase-2) |
|  | L20688 | tho GDP-dissociation inhibitor protein 2 (Ly-GDI) |
|  | L26081 | semaphorin III |
|  | L34056 | CADHERIN-11 (OSTEOBLAST-CADHERIN) (OB-CADHERIN) |
|  | L34057; [L33477] | CADHERIN-12 (BR-CADHERIN) (N-CADHERIN 2) (CADHERIN, NEURAL TYPE, 2) |


|  | GenBank \# | CELL INTERACTION (Gene Names) |
| :---: | :---: | :---: |
|  | L34058; [U59289; U59288] | CADHERIN-13 T-CADHERIN PRECURSOR (TRUNCATED-CADHERIN) (H-CADHERIN) (HEART-CADHERIN) |
|  | L34059 | CADHERIN-4 RETINAL-CADHERIN PRECURSOR (R-CADHERIN) (R-CAD) |
|  | $\cdots$ … ${ }^{\text {a }}$ - ... 34060 | CADTEAIN-8 |
|  | M23410 | PLAKOGLOBIN (DESMOPLAKIN III) |
|  | M94151 | ALPHA-CATENIN RELATED PROTEIN (CATENIN ALPHA-2) |
|  | U24152 | SERINE/THREONINE-PROTEIN KINASE PAK-ALPHA (EC 2.7.1.-) (P65-PAK) (P21ACTIVATED KINASE) (ALPHA-PAK) |
|  | U24153 | P21-activated protein kinase (Pak2) |
|  | U33920 | semaphorin V |
|  | U43318 | frizzled 5 |
|  | X04429 | PLASMINOGEN ACTIVATOR INHIBITOR-1 PRECURSOR, ENDOTHELIAL (PAI-1) |
|  | X13916 | LOW-DENSITY LIPOPROTEIN RECEPTOR-RELATED PROTEIN 1 PRECURSOR (LRP) (ALPHA-2-MACROGLOBULIN RECEPTOR) (A2MR) |
|  | X14787 | THROMBOSPONDIN 1 PRECURSOR |
|  | L40027 | glycogen synthase kinase 3 |
| $\pm$ | X54412 | collagen type IX alpha-1 |
| 9 | X56654 | desmoglein type 1 |
|  | $\times 56807$ | DSC2 mRNA for desmocollins type 2a and 2b |
|  | X61587 | rhoG |
|  | X63629 | CADHERIN-3 PLACENTAL-CADHERIN PRECURSOR (P-CADHERIN) |
|  | X69550 | rho GDP-dissociation Inhibitor 1 |
|  | X75308 | MMP-13 (collagenase-3) |
|  | $\times 78565$ | TENASCIN-C |
|  | $\begin{aligned} & \times 79981 ; \\ & \times 59796] \end{aligned}$ | CADHERIN-5 VASCULAR ENDOTHELIAL-CADHERIN PRECURSOR (VE-CADHERIN) (7B4 ANTIGEN) (CD144 ANTIGEN). |
|  | M11313 | ALPHA-2-MACROGLOBULIN PRECURSOR (ALPHA-2-M) |
|  | X95282 | Rho8 protein |
|  | X95456 | Rho7 protein |
|  | Y07923 | Rho6 protein |
|  | 213009 | CADHERIN-1(E-CADHERIN) (UVOMORULIN) (CAM 120/80) |
|  | 215009 | laminin |
|  | 248482 | MMP-15 (MT2-MMP) |
|  | AB000220 | semaphorin E |
|  | - AFOO3522 | Della |


|  | GenBank \＃ | CELL INTERACTION（Gene Names） |
| :---: | :---: | :---: |
|  | D85815 | rhoHP1 |
|  | AF000974 | Zyxin related protein ZRP－1 |
|  | U29343 | HYALURONAN RECEPTOR（RHAMM） |
|  | M24795 | PLATELET GLYCOPROTEIN IV（GPIV）（GPIIB）（CD36 ANTIGEN）（PAS IV）（PAS－4 PROTEIN） |
|  | U72661 | NINJURIN－1 |
|  | U76456 | TIMP．4 |
|  | U82532 | GDI－dissociation inhibitor RhoGDIgammma |
|  | X92521 | MMP－19 |
|  | Y07604 | nm23－H4；NUCLEOSIDE－DIPHOSPHATE KINASE（EC 2．7．4．6）（NUCLEOSIDE 5＇－ DIPHOSPHATE PHOSPHOTRANSFERASE）（NDK）． |
|  | Y11306 | beta catenin／TCF． 4 |
|  | U38276 | SEMAPHORIN－1 |
|  | U94354 | lunatic fringe |
|  | U02570 | CDC42 GTPase－activating protein |
|  | X05199 | PLASMINOGEN PRECURSOR（EC 3．4．21．7） |
| $\underline{1}$ | $\times 05231$ | MMP－1（collagenase－1） |
| $\cdots$ | X53795 | CD82 ANTIGEN（INDUCIBLE MEMBRANE PROTEIN R2）（C33 ANTIGEN）（IA4） （METASTASIS SUPPRESSOR KANGAI 1）（SUPPRESSOR OF TUMORIGENICITY－6）． |
|  | L38517 | indian hedgehog protein（ 1 HH ） |
|  | M31470 | ras－like protein TC10 |
|  | M34189 | integrin betal |
|  | $\begin{aligned} & \text { X83929; } \\ & {[D 17427]} \end{aligned}$ | desmocollin type $3+$ desmocollin type 4 |
|  | L23808 | MMP－12（metalloelastase） |
|  | L25081 | rhoC（H9）；SMALL GTPase（rhoC） |
|  | $\begin{aligned} & \text { M29870; } \\ & \text { [M31467] } \end{aligned}$ | RAS－RELATED C3 BOTULINUM TOXIN SUBSTRATE 1 （P21－RAC1）（RAS－LIKE PROTEIN TC25） |
|  | $\begin{aligned} & \text { M64595; } \\ & {[\mathrm{M} 29871]} \end{aligned}$ | RAS－RELATED C3 BOTULINUM TOXIN SUBSTRATE 1 （P21－RAC2） |
|  | X05232 | MMP－3（stromelysin－1） |
|  | $\times 06820$ | rhoB |
|  | $\begin{aligned} & \text { X07820, } \\ & {[M 30461]} \end{aligned}$ | MMP－10（stromelysin－2） |
|  | $\times 72925$ | desmocollin type 1 |




## C'ytokine and C'ytokine Receptor Array

In the cytokine and cytokine receptor array according to the subject invention, all of the unique polynucleotide probe compositions correspond to genes that express cytokines or cytokine receptors. In a specific cytokine and cytokine receptor array of interest, the spots are as provided in Table 8.

|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | M29696 | INTERLEUKIN-7 RECEPTOR ALPHA CHAIN |  |
|  | X01992 | INTERFERON GAMMA |  |
|  | J04156 | INTERLEUKIN-7 |  |
|  | X01057 | INTERLEUKIN-2 RECEPTOR ALPHA CHAIN |  |
|  | A14844 | INTERLEUKIN-2 |  |
|  | M29366 | PROTEIN-TYROSINE KINASE RECEPTOR ERBB-3 [Epidermal growth factor receptor (avian erythroblastic leukemia viral (v-erb-b) oncogene homolog)ll |  |
|  | X04434 | INSULIN-LIKE GROWTH FACTOR I RECEPTOR |  |
|  | M29645 | INSULIN-LIKE GROWTH FACTOR II [Somatomedin A] |  |
|  | X03663 | MACROPHAGE COLONY STIMULATING FACTOR I RECEPTOR [c-ims proto-oncogene] |  |
|  | $\begin{aligned} & \begin{array}{l} \text { M32315; } \\ \text { M55994 } \end{array} \end{aligned}$ | TUMOR NECROSIS FACTOR RECEPTOR 2 PRECURSOR (TUMOR NECROSIS FACTOR BINDING PROTEIN 2) (TBPII) (P80) (TNF-R2) (P75) (CD120B) (TNFR2) (TNFBR). |  |
|  | $\begin{aligned} & \mathrm{X02811;} \\ & {[\mathrm{X} 02744 ;} \\ & \text { M12783 } \end{aligned}$ | PLATELET-DERIVED GROWTH FACTOR, B CHAIN PRECURSOR (PDGF B-CHAIN) (PDGF-2) (BACAPLERMIN) (C-SIS) |  |
|  | $\times 02851$ | INTERLEUKIN-1 ALPHA |  |
|  | K02770 | INTERLEUKIN IL-1BETA |  |
| $\frac{1}{N}$ | $\begin{aligned} & \text { M14743; } \\ & {[\text { M17115] }} \end{aligned}$ | INTERLEUKIN-3 PRECURSOR (IL-3) (MULTIPOTENTIAL COLONY-STIMULATING FACTOR) (HEMATOPOIETIC GROWTH FACTOR) (P-CELL STIMULATING FACTOR) (MAST-CELL GROWTH FACTOR) (MCGF) (IL3). |  |
| 1 | M13982 | INTERLEUKIN-4 |  |
|  | X04602; <br> [M14584] | INTERLEUKIN-6 PRECURSOR (IL-6) (B-CELL STIMULATORY FACTOR 2) (BSF-2) (INTERFERON BETA-2) (HYBRIDOMA GROWTH FACTOR). |  |
|  | X01394 | TUMOR NECROSIS FACTOR [TNFa]. |  |
|  | D12614 | LYMPHOTOXIN-ALPHA [formerly fumor necrosis factor beta (TNF-beta)] |  |
|  | M20566 | INTERLEUKIN-6 RECEPTOR ALPHA CHAIN |  |
|  | $\left[\begin{array}{l} x 04688 ; \\ (J 03478) \end{array}\right.$ |  FACTOR) (EOSINOPHIL DIFFERENTIATION FACTOR) |  |
|  | M28622 | INTERFERON BETA |  |
|  | M11220 | GRANULOCYTE-MACROPHAGE COLONY-STIMULATING FACTOR [GM-CSF] |  |
|  | K03222 | TRANSFORMING GROWTH FACTOR-ALPHA |  |
|  | $\begin{aligned} & \text { J00209; } \\ & \text { jJ00207] } \end{aligned}$ | LEUKOCYTE INTERFERON ALPHA |  |
|  | $\times 02812$ | TRANSFORMING GROWTH FACTOR BETA [1] |  |
|  | X03438 | GRANULOOCYẎE COLONY-STIMULATING FACTOR [G-CSF] |  |
|  | M19154 |  |  |
|  | $\times 04571$ | E'PIDERMAL GROWTHFACTOR KIDNEY [EGF] |  |
|  | J03171 | HUIFN-ALPHA -REC JINTERFERON ALPHA-BETA RECEPTOR ALPHA CHAIN] |  |
|  | M57627 | INTERLEUKIN-10 |  |
|  | M26062 | İNTERLEUKIN-2 RECEPTOR BETA CHAIN |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | M74782 | INTERLEUKIN-3 RECEPTOR ALPHA CHAIN |  |
|  | $\times 1$. | INTERLEUKIN-4 RECEPTOR ALPHA CHAIN |  |
|  | M75914 | INTERLEUKIN-5 RECEPTOR ALPHA CHAIN |  |
|  | X 77722 | INTEAFEERON ALPHA-BETA RECEPTOR BETA CHAIN |  |
|  | X7275 5 | GAMMA INTERFERON INDUCED MONOKINE [Humig] |  |
|  | D11086 | CYTOKINE RECEPTOR COMMON GAMMA CHAIN [Interleukin 2 receptor gamma chain] |  |
|  | M20132 | ANDROGEN RECEPTOR |  |
|  | M73238 | CILIARY NEUROTROPHIC FACTOR RECEPTOR ALPHA |  |
|  | J03143 | INTERFERON-GAMMA RECEPTOR ALPHA CHAIN |  |
|  | M60459 | ERYTHROPROTEIN RECEPTOR |  |
|  | L00587 | CALCITONIN RECEPTOR |  |
|  | M62424 | THROMBIN RECEPTOR [Coagulation factor II (thrombin) receptor] |  |
|  | L07594 | TRANSFORMING GROWTH FACTOR-BETA TYPE III RECEPTOR |  |
|  | M84747 | INTERLEUKIN-9 RECEPTOR |  |
|  | U00672 | INTERLEUKIN-10 RECEPTOR |  |
|  | M14764 | LOW-AFFINITY NERVE GROWTH FACTOR RECEPTOR |  |
|  | $\begin{aligned} & X 60957 \\ & (S 89716) \end{aligned}$ | TYROSINE-PROTEIN KINASE RECEPTOR TIE-1 PRECURSOR (EC 2.7.1.112). |  |
| N | $\begin{aligned} & \times 68203 ; \\ & (\times 69878 ; \\ & \text { U43143) } \end{aligned}$ | V̄ĀS̄C̄ULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR 3 PRECURSOR (EC 2.7.1.112) (VEGFR-3) (TYROSINE-PROTEIN KINASE RECEPTOR FLT4, CLASS III). |  |
|  | M16552 | THROMBOMODULIN |  |
|  | M87290 | ANGIOTENSIN II RECEPTOR TYPE-1A |  |
|  | M83941 | TYROOSINE-PROTEIN KINASE RECEPTOR ETK1 |  |
|  | M76673 | FMLP-RELATED RECEPTOR I |  |
|  | M97675 | TRANSMEMBRANE RECEPTOR ROR1 |  |
|  | $\begin{aligned} & L 04947 ; \\ & {[\times 61656]} \end{aligned}$ | VASCULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR 2 PRECURSOR (EC 2.7.1.112) (VEGFR-2) (KDR) (KINASE INSERT DOMAIN RECEPTOR) (FRAGMENT) |  |
|  | M91196 | INTERFERON CONSENSUS SEQUENCE BINDING PROTEIN [DNA-binding protein] |  |
|  | $\times 75208$ | TYROSINE-PROTEIN KINASE RECEPTOR EPH-3 |  |
|  | U05012 | trk-C |  |
|  | X74764 | TYROSINE-PROTEIN KINASE CAK [Tyrosine kinase, receptor TKT] |  |
|  | K03193;  <br> IX00588;  <br> X00663;  <br> U48722  <br> D10202  | EPIDERMAL GROWTH FACTOR RECEPTOR PRECURSOR (EC 2.7.1.112). (EGFR) (ERBB1) <br> PLATELET AC TIVATING FACTOR RECEPTOR | ..... .- |
|  | M18391 | TYROSINE-PROTEIN KINASE RECEPTOR EPH |  |
|  | A09781 | INTERFERON-GAMMA RECEPTOR |  |
|  | U12140 | TYROSINE KINASE RECEPTOR TRK-B |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | M86492 | GLIȦ MATURATION FACTOR BETA |  |
|  | L07868 | ERBB4 [EPIDERMAL GROWTH FACTOR RECEPTOR] |  |
|  | M27492 | INTERLEUKIN-1 RECEPTOR TYPE I |  |
|  | M 33294 | TUMOR NECROSIS FACTOR RECEPTOR 1 |  |
|  | M37435 | MACROPHAGE COLONY STIMULATING FACTOR-1 [M-CSF] |  |
|  | M11730 | ERBB-2 RECEPTOR PROTEIN-TYROSINE KINASE |  |
|  | D10923 | HM74 [PROBABLE G PROTEIN-COUPLED RECEPTOR HM74] |  |
|  | D10924 | HM89 [PROBABLE G PROTEIN-COUPLED RECEPTOR LCR1 HOMOLOG] |  |
|  | D10925 | HM145 [C-C CHEMOKINE RECEPTOR TYPE 1] |  |
|  | 014012 | HEPATOCYTE GROWTH FACTOR ACTIVATOR |  |
|  | D16431 | HEPTOMA-DERIVED GROWTH FACTOR |  |
|  | $\begin{aligned} & \text { D30751; } \\ & \text { (M22490) } \end{aligned}$ | BONE MORPHOGENETIC PROTEIN 4 (BMP-2B) |  |
|  | j03358 | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE FER |  |
|  | 504130 | MACROPHAGE INFLAMMATORY PROTEIN 1-BETA [Activation (Act-2)] |  |
|  | J05081 | ENDOTHELIN-3 |  |
|  | L06139 | TYROSINE-PROTEIN KINASE RECEPTOR TIE-2 PRECURSOR (EC 2.7.1.112) (TYROSINE PROTEIN KINASE RECEPTOR TEK) (P140 TEK) (TUNICA INTERNA ENDOTHELIAL CELL KINASE). |  |
|  | L06622 | ENDOTHELIN-1 RECEPTOR [EDNRA] |  |
|  | L06623 | ENDOTHELIN B RECEPTOR [EDNRB] |  |
|  | L06801 | INTERLEUKIN-13 |  |
|  | L07414 | CD40 LIGAND |  |
|  | L08096 | CD27 LIGAND [CD70 anligen] |  |
|  | L08187 | CILIARY NEUROTROPHIC FACTOR RECEPTOR ALPHA [cytokine receptor EB13] |  |
|  | L09753 | CD30 |  |
|  | $\begin{aligned} & \text { L12260; } \\ & \text { U02326; } \\ & \text { M94165 } \\ & \hline \end{aligned}$ | RECOMBINANT GLIAL GROWTH FACTOR + NEU DIFFERENTIATION FACTOR + HEREGULIN |  |
|  | L12261 | HEREGULIN ALPHA [Recombinant glial growth factor] |  |
|  | L15344 | INTERLEUKIN IL-14 |  |
|  | $\begin{aligned} & \text { L36052; } \\ & \text { (L36051; } \\ & \text { U1 1025) } \end{aligned}$ | THROMBOPOIETIN PRECURSOR (MEGAKARYOCYTE COLONY STIMULATING FACTOR) (C-MPL LIGAND) (ML) (MEGAKARYOCYTE GROWTH AND DEVELOPMENT FACTOR) (MGDF) (THPO) |  |
|  | M10051 | INSULIN RECEPTOR |  |
|  | M21121 | RANTES PROTEIN T-CELL SPECIFIC |  |
|  | M21574 | PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA |  |
|  | M21616 | PLATELET-DERIVED GROWTH FACTOR RECEPTOR BETA |  |
|  | $\begin{aligned} & \text { M22488 } \\ & \text { (U50330] } \end{aligned}$ | BONE MORPHOGENETIC PROTEIN 1 (procollagen C-proteinase) (pCP-2) |  |
|  | M22489 | BONE MORPHOGENETIC PROTEIN 2A |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | M22491 | BONE MORPHOGENETIC PROTEIN 3 |  |
|  | M23452 | MÁCROPHAGE INFLAMMATORY PROTEIN 1-ALPHA [GOS19-1] |  |
|  | M24545 | MONOCYTE CHEMOTACTIC PROTEIN 1 |  |
|  | M25667 | NEUROMODULIN [Neuronal growth protein 43 (GAP-43)] |  |
|  | M27288 | ONCOSTATIN M |  |
|  | M30704 | AMPHIREGULIN [schwannoma-derived growth factor] |  |
|  | M31145 | INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 1 |  |
|  | M31165 | TUMOR NECROSIS FACTOR-INDUCIBLE PROTEIN TSG-6 |  |
|  | M32977; <br> (M27281) | VASCULAR ENDOTHELIAL GROWTH FACTOR PRECURSOR (VEGF) (VASCULAR PERMEABILITY FACTOR) (VPF). |  |
|  | M35410 | IGFBP-2 [INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 2 |  |
|  | M36717 | PLACENTAL RIBONUCLEASE INHIBITOR [Ribonuclease/angiogenin inhibitor RAI] |  |
|  | M37722; <br> (X66945; <br> M63887: <br> M63888; <br> M63889;M3418 <br> 6; M34641] | BASIC FIBROBLAST GROWTH FACTOR RECEPTOR 1 PRECURSOR (BFGF-R) (EC 2.7.1.112) (FMS-LIKE TYROSINE KINASE-2) (C-FGR) (FGFR1) (FLG) (FGFBR) (FLT2). (HBGF-R-ALPHA-A1) (HBGF-R-ALPHA-A2) (HBGF-R-ALPHA-A3) + FGFR SECRETED FORM (M34188) |  |
|  | $\text { M } 57230$ | INTERLEUKIN-6 RECEPTOR BETA CHAIN [membrane glycoprotein gp130] |  |
| $\frac{1}{N}$ | $\begin{aligned} & \text { M57399; } \\ & \text { [X52946; } \\ & \mathrm{D} 90226] \end{aligned}$ | PLEIOTROPHIN PRECURSOR (PTN) (HEPARIN-BINDING GROWTH-ASSOCIATED MOLECULE) (HB-GAM) (HEPARIN-BINDING GROWTH FACTOR 8) (HBGF-8) (OSTEOBLAST SPECIFIC FACTOR 1) (OSF-1) (HEPARIN-BINDING NEURITE OUTGROWTH PROMOTING FACTOR 1) (HBNF-1). |  |
|  | M57502 | TLYMPHOCYTE-SECRETED PROTEIN I-309 |  |
|  | M57765 | INTERLEUKIN-11 [adipogenesis inhibitory factor] |  |
|  | M59818 | GRANULOCYTE COLONY STIMULATING FACTOR RECEPTOR |  |
|  | M59964 | STEM CELL FACTOR (C-KIT LIGAND) |  |
|  | M60278 | HEPARIN-BINDING EGF-LIKE GROWTH FACTOR [DIPHTHERIA TOXIN RECEPTOR] |  |
|  | M60718 | HEPĀTOCYTE GROWTH FACTOR PRECURSOR (SCATTER FACTOR) (SF) (HEPATOPOEITIN A). |  |
|  | M60828 | F̈GF-7: KERATINOCYTE GROWTH FACTOR PRECURSOR (KGF) (FIBROBLAST GROWTH FACTOR-7) (HBGF-7). |  |
|  | M61176 | BRAIN-DERIVED NEUROTROPHIC FACTOR |  |
|  | M62302 | GDF-1 [GROWTH/DIFFERENTIATION FACTOR 1] |  |
|  | M 62505 | CSA ANAPHYLATOXIN CHEMOTACTIC RECEPTOR |  |
|  | M65199 | ENDOTHELIN-2 |  |
|  | M 65290 | INTERLEUKIN-12 BETA CHAIN [Natural killer cell stimulatory factor, p40] |  |
|  | M65291 | INTERLEUKIN-12 ALPHA CHAIN [Natural killer cell stimulatory factor, p 35 ] |  |
|  | M67454 | F-ASL RECEPTOR [Fas antigen, APO.1 antigen] |  |
|  | M 68932 | INTERLEUKIN-8 RECEPTOR (ALFA, HIGH AFFINITY) |  |
|  | M73482 | NEUROMEDIN-B RECEPTOR |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | M74178 | HEPATOCYTE GROWTH FACTOR-LIKE (macrophage-stimulating protein (MST1)) |  |
|  | M76125 | AXL (TYROSINE-PROTEIN KINASE RECEPTOR UFO) |  |
|  | M92381 | THYMOSIN BETA-10 |  |
|  | M92934 | CONNECTIVE TISSUE GROWTH FACTOR |  |
|  | $\begin{aligned} & \text { M96956 } \\ & {[\text { M96955] }} \end{aligned}$ | T̄DGF1 (TERATOCARCINOMA-DERIVED GROWTH FACTOR 1) (EPIDERMAL GROWTH FACTOR-LIKE CRIPTO PROTEIN CR1) (CRIPTO-1 GROWTH FACTOR) (CRGF) + TDGF2 (TERATOCARCINOMA-DERIVED GROWTH FACTOR 2) (EPIDERMAL GROWTH FACTOR-LIKE CRIPTO PROTEIN CR3) (CRIPTO-3 GROWTH |  |
|  | 559184 | TYROSINE.PROTEIN KINASE RYK [RYK receptor-like tyrosine kinase] |  |
|  | $\begin{aligned} & \text { U01134; } \\ & {[\times 51602]} \end{aligned}$ | VASCULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR 1 PRECURSOR (EC 2.7.1.112) (VEGFR-1) (TYROSINE-PROTEIN KINASE RECEPTOR FLT) (FLT-1) (SFLT) |  |
|  | U02687 | FL CYTOKINE RECEPTOR PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR FLT3) (STEM CELL TYROSINE KINASE 1) (STK-1) (CD135 ANTIGEN). |  |
|  | 003187 | İNTERLEUKIN-12 RECEPTOR |  |
|  | U03882 | C-C CHEMOKINE RECEPTOR [Monocyte chemoattractant protein 1 receptor (MCP-1RA) alternatively spliced) |  |
|  | 003905 | C.C CHEMOKINE RECEPTOR [Monocyte chemoattractant protein 1 receptor (MCP-1RB) allernatively spliced] |  |
| $\frac{1}{N}$ | U04806; (U03858) | SL CYTOKINE PRECURSOR (FLT3/FLK2 LIGAND). |  |
| Y | U10117 | ENDOTHELIAL-MONOCYTE ACTIVATING POLYPEPTIDE II |  |
|  | $\begin{aligned} & \text { U11814; } \\ & \text { [M80634; } \\ & \text { X52832; } \\ & \text { M35718; } \\ & \text { M87771; } \\ & M 87772 \text { ] } \end{aligned}$ | FIBROBLAST GROWTH FACTOR RECEPTOR 2 PRECURSOR (FGFR-2) (EC 2.7.1.112) (KERATINOCYTE GROWTH FACTOR RECEPTOR) (FGFR2) (BEK) (BFR-1) (KSAM-1) + KSAM; K-SAM-III; K-SAM-IV |  |
|  | U14407 | INTERLEUKIN-15 |  |
|  | U14722 | ACTIVIN TYPE I RECEPTOR |  |
|  | U43142 | VASCULAR ENDOTHELIAL GROWTH FACTOR C PRECURSOR (VEGF-C) (VASCULAR ENDOTHELIAL GROWTH FACTOR RELATED PROTEIN) (VRP) (FLT4 LIGAND). |  |
|  | $\times 06182$ | C-KIT PROTO-ONCOGENE [masUstem cell growth factor receptor] |  |
|  | $\times 1$ | CALGRANULIN (B) [MRP-14 (calcium binding protein in macrophages,MIF-related)] |  |
|  | X06234 | CALGRANULIN (A) [MRP-8 (calcium binding protein in macrophages,MIF-related)] |  |
|  | X06374 | PLATELET-DERIVED GROWTH FACTOR (A CHAIN) [PDGF-A] |  |
|  | $\times 13967$ | LEUKAEMIA INHIBITORY FACTOR [cholinergic differentiation factor] |  |
|  | X17543 | INTERLEUKIN-9 |  |
|  | $\times 17648$ | GRANULOC YTE-MACROPHAGE COLONY-STIMULATING FACTOR RECEPTOR ALPHA CHAIN [hGM-CSF-R] |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | X51943: (M13361: X65778 | HEPARIN-BINDING GROWTH FACTOR 1 PRECURSOR (HBGF-1) (ACIDIC FIBROBLAST GROWTH FACTOR) (AFGF) (BETA-ENDOTHELIAL CELL GROWTH FACTOR) (ECGFBETA). |  |
|  | x53655: | N̄T- 3 (NEUROTROPHIN-3 PRECURSOR) (NEUROTROPHIC FACTOR) (HDNF) (NERVE GROWTH FACTOR 2) (NGF-2) |  |
|  | X53799 | MACROPHAGE INFLAMMATORY PROTEIN-2-ALPHA [MIP2alpha] |  |
|  | $\times 54936$ | PLACENTA GROWTH FACTORS 1 AND 2 PRECURSOR (PLGF-1/ PLGF-2). |  |
|  | X59770 | INTERLEUKIN-1 RECEPTOR TYPE II |  |
|  | X60592 | CDW40; NERVE GROWTH FACTOR RECEPTOR-RELATED B-LYMPHOCYTE ACTIVATION MOLECULE |  |
|  | x72304 | CORTICOTROPIN RELEASING FACTOR RECEPTOR |  |
|  | X78686 | NEUTROPHIL ACTIVATING PROTEIN ENA-78 |  |
|  | X79929 | OX40 LIGAND [9p34] |  |
|  | Y00787 | INTERLEUKIN-8 [monocyte-derived neutrophil chemolactic factor MDNCF] |  |
|  | 270519 | FAS/APO 1 |  |
|  | D17517 | TYROSINE-PROTEIN KINASE RECEPTOR UFO [sky] |  |
|  | J03241 | TRANSFORMING GROWTH FACTOR (BETA 3) |  |
|  | J03634 | INHIBIN BETA (A CHAIN) [activin A, activin AB alpha polypeptide; erythroid differentiation protein mRNA (EDF)] |  |
|  | L32976 | PROTEIN KINASE MLK-3 [MIXED LINEAGE KINASE 1] |  |
| - | L35233 | AUTOCRINE MOTLLITY FACTOR RECEPTOR [AMFR] |  |
|  | M31213: [M57464] | PROTO-ONCOGENE TYROSINE-PROTEIN KINASE RECEPTOR RET PRECURSOR (EC 2.7.1.112) (C-RET). (Papillary thyroid carcinoma-encoded protein] |  |
|  | M95489 | FOLLICLE STIMULATING HORMONE RECEPTOR |  |
|  | U05875 | INTERFERON-GAMMA RECEPTOR BETA CHAIN $\quad$ IInterferon gamma receplor accessory facior-1 (AF-1)] |  |
|  | $\begin{aligned} & 15979 ; \\ & 1212172] \\ & \hline \end{aligned}$ | DELTA-LIKE PROTEIN PRECURSOR (CONTAINS: FETAL ANTIGEN 1) (FA1) (DLK) + ADRENAL SPECIFIC 30 kd PROTEIN GB: X17544 |  |
|  | $\times 03541$ | HiGh Af IINITY NERVE GROWTH FACTOR RECEPTOR PRECURSOR (EC 2.7.1.112) (TRK1 TRANSFORMING TYROSINE KINASE PROTEIN) (P140-TRKA) + trk-T3 (P68 TRKT3 ONCOPROTEIN) |  |
|  | $\times 15218$ | SKI ONCOGENE |  |
|  | $\times 15219$ | SKI-RELATED ONCOGENE SNON |  |
|  | X74979 | TYROSINE-PROTEIN KINASE CAK [EDDR1; TRKE] |  |
|  | A06925 | RELAXIN H2 |  |
|  | 010232 | RENIN-BINDING PROTEIN |  |
|  | M13981 | INHIBIN ALP PHA CHAIN |  |
|  | M31159; | IGFBP3 (GROWTH HORMONE-DEPENDENT INSULIN-LIKE GROWTH FACTOR-BINDING |  |
|  | U06863 | FOLLISTATIN-RELATED PROTEIN |  |
|  | S85655 | PROHIBITIN |  |


| GenBank \# | Gene Name |  |
| :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { D38122; } \\ & \text { [U08137] } \end{aligned}$ | FAS ANTIGEN LIGAND (APOPTOSIS ANTIGEN LIGAND) (APTL) (APT1LG1) (FASL). |  |
| L11015 | LYMPHOTOXIN-BETA |  |
| U57059 | FAS ANTIGEN LIGAND [TNF-related apoptosis inducing ligand TRAIL; Apo-2 ligand] |  |
| X14454 | INTERFERON REGULATORY FACTOR [Interferon regulatory factor 1] |  |
| $\begin{aligned} & \text { Y09392; } \\ & \text { (U75380;U7461 } \\ & 1 ; \text { U83597) } \end{aligned}$ | WSL-LR, WSL-S1, WSL-S2 + TRAMP (Apo-3) (DDR3) |  |
| M27544 | INSULIN-LIKE GROWTH FACTOR IA |  |
| M86528 | NEUROTROPHIN-4 |  |
| M86528; <br> S41541: <br> [S41540; <br> S41522 | NT-4 (NT-5) + NT-6 |  |
| U14187 | RECEPTOR TYROSINE KINASE LIGAND LERK-3 (EPLG3) |  |
| U14188 | RECEPTOR TYROSINE KINASE LIGAND LERK-4 (EPLG4) |  |
| U32659 | INTERLEUKIN-17 |  |
| U33635 | HIGH AFFINITY NERVE GROWTH FACTOR RECEPTOR [colon carcinoma kinase-4 (CCK4)] |  |
| U68162 | THROMBOPOEITIN RECEPTOR |  |
| A25270 | IFN-GAMMA ANTAGONIST CYTOKINE |  |
| A03911 | NEURITE PROMOTING FACTOR(NEXIN), glia derived |  |
| D49493 | BONE MORPHOGENETIC PROTEIN 3B |  |
| $\begin{aligned} & \hline \text { D49742; } \\ & {[\text { S83182 }]} \end{aligned}$ | HGF ACTIVATOR LIKE |  |
| L17075 | TGF-b superfamily receptor type I (ALK-1) (SRK3) |  |
| L03840 | FGFR4 |  |
| L19063 | GDNF |  |
| L37882 | frizzled |  |
| L20861 | Wnt-5a |  |
| M62403 | IGFBP4 |  |
| M65062 | IGFBP5 |  |
| M73980 | Notch1 |  |
| M97016 | BONE MORPHOGENETIC PROTEIN 8 (OSTEOGENIC PROTEIN 2) |  |
| M99437 | notch group protein (N) |  |
| U43318 | frizzled 5 |  |
| $\times 07876$ | WNT2 OR IRP |  |
| A26792 | CNTF, ISOFORM B AND C |  |
| L42379 | BPGF. 1 |  |
| 271621 | Wnt-13 |  |
| M21626 | T CELL RECEPTOR VARIABLE REGION |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | M25639 | MIF |  |
|  | U82169 | frizzled homolog (FZD3) |  |
|  | U83508 | angiopoietin-1 |  |
|  | U84401 | smoothened |  |
|  | U90875 | cytotoxic ligand TRAIL receptor |  |
|  | U95299 | Notch4 |  |
|  | X91940 | WNT-8B |  |
|  | X97057 | WNT-10B |  |
|  | AF003521 | Jagged 2 |  |
|  | AF028593 | Jagged 1 |  |
|  | U77493 | Notch2 |  |
|  | U94352 | manic fringe |  |
|  | U94354 | lunatic fringe |  |
|  | M27968 | FGF2; HEPARIN-BINDING GROWTH FACTOR 2 PRECURSOR (PROSTATROPIN). (HBGF 2) (BASIC FIBROBLAST GROWTH FACTOR) (BFGF) (PROSTATROPIN) |  |
|  | L38518 | sonic hedgehog (SHH) |  |
|  | M60314 | BONE MORPHOGENETIC PROTEIN 5 |  |
|  | M60315 | BONE MORPHOGENETIC PROTEIN 6 |  |
|  | M60316 | BONE MORPHOGENETIC PROTEIN 7 (OSTEOGENIC PROTEIN 1) |  |
|  | $\begin{aligned} & \text { D13365; } \\ & \text { M93311] } \end{aligned}$ | GROWTH INHIBITORY FACTOR (METALLOTHIONEIN-III) (MT-III) |  |
|  | U46010 | HGF AGONIST/ANTAGOINST |  |
|  | L36034 | SDF 1A (pre-B cell stimulating factor homologue) |  |
|  | M15530 | BCGF1 (B-cell growth factor) |  |
|  | $\begin{aligned} & \overline{M 58051 ;} \\ & \times 58255 \end{aligned}$ | FGFR3 (FLG-2) |  |
|  | M77227 | COMPETITIVE HEPATOCYTE GROWTH FACTOR ANTAGONIST. AN ALTERNATIVE TRANSCRIPT OF THE HEPATOCYTE GROWTH FACTOR PRECURSOR (SCATTER FACTOR) (SF) (HEPATOPOEITIN A) |  |
|  | $\begin{aligned} & \text { U24163; } \\ & \text { [U91903; } \\ & \text { U68057] } \end{aligned}$ | frizzled-related FrzB (Frizz) (frezzled (fre)) |  |
|  | $\begin{aligned} & \text { U28811; } \\ & \text { (U64791) } \end{aligned}$ | CYSTEINE-RICH FIBROBLAST GROWTH FACTOR RECEPTOR [Golgi membrane sialoglycoprotein MG160 (GLG1)] |  |
| - . .. | $\begin{aligned} & U 48801 \text {; } \\ & \text { [U43368 } \\ & \times 02492 \end{aligned}$ | VASCULAR ENDOTHELIAL GROWTH FACTOR B PRECURSOR (VEGF-B) + VEGF RELATED FACTOR ISOFORM VRF186 PRECURSOR L̈ËUKOCYTE INTERFERON-INDUCIBLE PEPTIDE |  |
|  | - $\times 1$ | trk-T3 (P68 TRK-T3 ONCOPROTEIN) |  |
|  | X14445 | FGF-3 : INT-2 PROTO-ONCOGENE PROTEIN PRECURSOR (FIBROBLAST GROWTH FACTOR-3)(HBGF-3). |  |
|  | M37825 | FGF-5; FIBROBLAST GROWTH FACTOR-5 PRECURSOR (HBGF-5). |  |


|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | AF022385 | apoptosis-related protein TFAR15 (TFAR15) |  |
|  | L20471 | extracellular matrix metalloproteinase inducer EMMPRIN |  |
|  | $\begin{aligned} & \text { M57730 } \\ & \text { M37476 } \end{aligned}$ | EPHRIN-A1 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 1) (LERK-1) (IMMEDIATE EARLY RESPONSE PROTEIN B61) (TUMOR NECROSIS FACTOR, ALPHA-INDUCED PROTEIN 4). |  |
|  | U07695 | EPHRIN TYPE-B RECEPTOR 4 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR HTK). |  |
|  | U09304 | EPHRIN-B1 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 2) (LERK-2) (ELK LIGAND PRECURSOR) (ELK-L). |  |
|  | U82938 | CD27BP (Siva) |  |
|  | U26403 | EPHRIN-AS PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 7) (LERK-7) (AL-1). |  |
|  | U66406 | EPHRIN-B3 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 8) (LERK-8) (EPH-RELATED RECEPTOR TRANSMEMBRANE LIGAND ELK-L3). |  |
|  | X95425 | EPHRIN TYPE-A RECEPTOR 5 PRECURSOR (EC 2.7.1.112) TYROSINE-PROTEIN KINASE RECEPTOR EHK-1) (EPH HOMOLOGY KINASE-1) (RECEPTOR PROTEIN. TYROSINE KINASE HEK7). |  |
|  | M62402 | IGFBP6 |  |
|  | AF016268 | death receptor 5 (DR5) |  |
| $\frac{1}{1}$ | AF017986 | secreted apoptosis related protein 1 |  |
| N | AF017988 | secreted apoptosis related protein 3 (SARP3) |  |
|  | L38734 | EPHRIN-B2 PRECURSOR (EPH-RELATED RECEPTOR TYROSINE KINASE LIGAND 5) (LERK-5) (HTK LIGAND) (HTK-L). |  |
|  | M63099 | INTERLEUKIN 1 RECEPTOR ANTAGONIST |  |
|  | L40636 | EPHRIN TYPE-B RECEPTOR 1 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR EPH-2) (NET). |  |
|  | L41939 | EPHRIN TYPE-B RECEPTOR 2 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN EPH- <br> 3) (DRT) |  |
|  | M16591 | TYROSINE-PROTEIN KINASE HCK (EC 2.7.1.112) (P59-HCK AND P60-HCK) (HEMOPOIETIC CELL KINASE). |  |
|  | $\begin{aligned} & \text { M59371 } \\ & \text { M36395 } \end{aligned}$ | EPHRIN TYPE-A RECEPTOR 2 PRECURSOR (EC 2.7.1.112) (TYROSINE-PROTEIN KINASE RECEPTOR ECK) (EPITHELIAL CELL KINASE). |  |
|  | D14838 | FGF-9; GLIA-ACTIVATING FACTOR PRECURSOR (GAF) (FIBROBLAST GROWTH FACTOR-9) (HBGF-9). |  |
|  | M77349 | BIGH3 |  |
|  | D25216 | IGFBP COMPLEX ACID LABILE CHAIN |  |
|  | U36223 | FGF-8; ANDROGEN-INDUCED GROWTH FACTOR PRECURSOR (AIGF) (HBGF-8) (FIBROBLAST GROWTH FACTOR-8) |  |
|  | U41745 | PDGF assoc. prolein |  |
|  | U43148 | patched homolog (PTC) |  |
|  | J02958 | MET |  |


|  | GenBank\# | Gene Name |  |
| :--- | :--- | :--- | :--- |
|  | U66197 | FHF-1 |  |
|  | $\times 52599$ | BETA NGF |  |
|  | retinoic acid receptor alpha [RETINOIC ACID RECEPTOR RXR-ALPHA (RXRA)I |  |  |
|  | X52773 | KGF-6; FIBROBLAST GROWTH FACTOR-6 PRECURSOR (HBGF-6) (HST-2). |  |
|  | X65923 | FAU |  |

## Cell Cycle Array

In the cell cycle array according to the subject invention, all of the unique polynucleotide probe compositions correspond to genes that are associated with the life cycle of a cell. In a specific cell cycle array of interest, the spots are as provided in Table 9.

|  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: |
|  | Z12020; [M92424] | MDM2 PROTEIN (P53-ASSOCIATED PROTEIN) + MDM2-A (GB: U33199) + MDM2-C (GB: U33201) |  |
|  | M14694: [M14695] | p53 |  |
|  | U18422 | DP2 (Humdp2) , dimerization partner ol E2F |  |
|  | L05624 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 1 (EC 2.7.1.-) (MAP KINASE KINASE 1) (MAPKK 1) (ERK ACTIVATOR KINASE 1) (MAPKKERK KINASE 1) (MEK1). |  |
|  | L07540 | ACTIVATOR 136 KD SUBUNIT (REPLICATION FACTOR C 36 KD SUBUNIT) (RFC36) |  |
|  | L07541 | ACTIVATOR 138 KD SUBUNIT (REPLICATION FACTOR C 38 KD SUBUNIT) (RFC38) |  |
|  | L20320 | CELL DIVISION PROTEIN KINASE 7 (EC 2.7.1.) (CDK-ACTIVATING KINASE) (CAK) (39 KD PROTEIN KINASE) (P39 MO15) (STK1) (CAK1). |  |
|  | L29511: [M96995] | GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2 ADAPTOR PROTEIN) (ASH PROTEIN). |  |
|  | L33264 | CDC2-RELATED KINASE PISSLRE |  |
|  | M63488 | REPLICATION PROTEIN A 70 KD DNA-BINDING SUBUNIT (RP-A) (RF-A) (REPLICATION FACTOR-A PROTEIN 1) (SINGLE STRANDED DNA-BINDING PROTEIN) |  |
|  | M74524 | HHR6A (YEAST RAD6 HOMOLOG) (UBIQITIN-CONJUGATING ENZYME) (UBCA) |  |
|  | M87338 | ACTIVATOR 140 KD SUBUNHT (REPLICATION FACTOR C 40 KD SUBUNIT) (RFC40) |  |
|  | M87339 | ACTIVATOR 137 KD SUBUNIT (REPLICATION FACTOR C 37 KD SUBUNIT) (RFC37) |  |
|  | U09579; [L25610) | C̄YCLIN-DEPENDENT KINASE INHIBITOR 1 (MELANOMA DIFFERENTIATION ASSOCIATED PROTEIN 6) (MDA-6) (P21) (CDK-INTERACTING PROTEIN 1) (CIP1) (WAF1) (CDKN1A) (CDKN1) (SD11) (PIC1) (CAP20) |  |
|  | M68520 | CELL DIVISION PROTEIN KINASE 2 (EC 2.7.1.-) (P33 PROTEIN KINASE) |  |
|  | M81933 | cdc25A; M-PHASE INDUCER PHOSPHATASE 1 (EC 3.1.3.48) |  |
|  | M92287 | CYCLIN D3 |  |
|  | M96684 | TRANSCRIPTIONAL ACTIVATOR PROTEIN PUR-ALPHA |  |
|  | $\times 51688$ | CYCLIN A |  |
|  | $\times 03484$ | RAF ONCOGENE |  |
|  | ×59798; (M64349) | CYCLIN D1 (CYCLIN PRAD1) (BCL-1 ONCOGENE) |  |
|  | D13639 [M90813] | CYCLIN D? |  |
|  | HT3218 [K00065] | SUPEROXIDE DISMUTASE [Superoxide dismutase 1 (Cu/Zn)] |  |
|  | D21235 | ŪV EXCISION REPAIR PROTEIN PROTEIN RAD23 [xeroderma pigmentosum group C repair complementing protein HHR23A] |  |
|  | U11791 [U12685] | CYCLINH |  |
|  | L26318 | STRESS-ACTIVATED PROTEIN KINASE JNK1 (EC 2.7.1.) (C-JUN N-TERMINAL KINASE 1) (JNK-46) |  |
|  | 127211 | CYCLIN-DEPENDENT KINASE 4 INHIBITOR A (CDK4I) (P16-INK4) (P16-INK4A) (MULTIPLE TUMOR SUPPRESSOR 1) (MTS1). (CDKN2A) |  |




|  |  | GenBank \# | Gene Name |  |
| :---: | :---: | :---: | :---: | :---: |
|  |  | $\times 80343$ | CYCLIN-DEPENDENT KINASE 5 ACTIVATOR PRECURSOR (CDK5 ACTIVATOR) (TAU PROTEIN KINASE II 23 KD SUBUNIT) (TPKII REGULATORY SUBUNIT) (P23) (P25) (P35). |  |
|  |  | $\times 85134$ | RBQ-3 |  |
|  |  | M15796; [J04718] | PCNA (CYCLIN) |  |
|  |  | AF001954 | growth inhibitor p33ING1 (ING1) |  |
|  |  | AF007111 | MDM2-like p53-binding protein (MDMX) |  |
|  |  | D89667 | C-myc binding protein |  |
|  |  | U66469 | p53-dependent cell growth regulator CGR19 |  |
|  |  | U77949 | CDC6-RELATED PROTEIN |  |
|  |  | U78876 | MEK KINASE 3 |  |
|  |  | Y11416 | p73, a monoallelically expressed p53-related protein |  |
|  |  | Y10479 | E2F-3 |  |
|  |  | U02570 | CDC42 GTPase-activating protein |  |
| - |  | L11285 | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 2 (EC 2.7.1.) (MAP KINASE KINASE 2) (MAPKK 2) (ERK ACTIVATOR KINASE 2) (MAPKJERK KINASE 2) (MEK2). |  |
| $\omega$ |  | M63167 | Akt1 (rac prolein kinase alpha, protein kinase B, c-Akt) |  |
| 1 |  | S57153; S57160 | RBP 1(RETINOBLASTOMA-BINDING PROTEIN) |  |
|  |  | U23435; U31089 | Abl interactor 2 (Abi-2) + Abl binding protein 3 (AbIBF3) [ArgBPIB] |  |
|  |  | M29870; [M31467] | RAS-RELATED C3 BOTULINUM TOXIN SUBSTRATE 1 (P21-RAC1) (RAS-LIKE PROTEIN TC25) |  |
|  |  | M96577 | E2F-1 pRB-binding protein |  |
|  |  | $\underline{\mathrm{U}} 25265$ | DUAL SPECIFICITY MITOGEN-ACTIVATED PROTEIN KINASE KINASE 5 (EC 2.7.1.-) (MAP KINASE KINASE 5) (MAPKK 5) (MAPKJERK KINASE 5). |  |
|  |  | $\times 66357$ | CELL DIVISION PROTEIN KINASE 3 (EC 2.7.1.-). |  |
|  |  | M74091 | CYCLIN C G1/S-SPECIFIC |  |
|  |  | M80629 | CDC2-RELATED PROTEIN KINASE CHED |  |
|  |  | S66431 | RBP2 retinoblastoma binding protein |  |
|  |  | 000001 | CDC27HS PROTEIN |  |
|  |  | U01038 | SERINETHREONINE-PROTEIN KINASE PLK (EC 2.7.1.-) (PLK-1) (STPK13) |  |
|  |  | D50310 | CYCLIN I |  |
|  |  | U18291 | CDC16HS. |  |
|  |  | U63131 | CDC37 HOMOLOG. |  |
|  |  | U69276 | GRB-IR / GRB10 |  |
|  |  | $\times 66358$ | SERINETHREONINE-PROTEIN KINASE KKIALRE |  |

## Other Representative Arrays

In a neuroarray according to the subject invention. all of the unique polynucleotide probe compositions will correspond to genes that are expressed in brain related tissues. Genes that are represented on the array are key genes, by which is meant that they have been reported to play primary roles in a variety of different biological processes in brain tissues. Genes of interest that may be represented on the array include: ion channel/transport proteins; receptors; cell cycle regulators; stress response proteins; apoptosis proteins; signal transduction proteins; transcriptional factors; growth factors/interleukins/hormones; oncogenes and tumor suppressors; cell surface/adhesion proteins; DNA synthesis/repair/recombination genes; and metabolic pathway enzymes.

In certain embodiments, of particular interest is an array having the following types of genes represented on its surface: nuclear proteins; endoplasmic reticulum proteins; golgi complex proteins; endosomal proteins; lysosomal proteins; peroxisomal proteins; mitochondrial proteins; cytoplasmic proteins; cytoskeletal proteins; plasma membrane proteins; post synaptic and dendritic proteins; axonal and nerve terminal proteins; secreted proteins, neuropeptides, hormones and growth factors; extracellular matrix proteins; astrocyte and oligodendroglial proteins; immune system proteins; developmentally regulated proteins; regionally regulated proteins; and disease related proteins.

Other representative arrays include: (1) rat arrays, in which each of the unique polynucleotide corresponds to a key rat gene; (2) blood arrays, in which each unique polynucleotide corresponds to a gene associated cells and tissues associated with the cardiovascular system; (3) rat stress arrays; and (4) mouse stress arrays, in which each unique polynucleotide corresponds to a gene associated with the stress response of murine cells.

## Methods of Using the Subject Arrays

The subject arrays find use in a variety of different applications in which one is interested in detecting the occurrence of one or more binding events between target nucleic acids and probes on the array and then relating the occurrence of the binding event(s) to the presence of a target(s) in a sample. In general, the device will be contacted with the sample suspected of containing the target under conditions sufficient for binding of any target
present in the sample to a complementary polynucleotide present on the array. Generally, the sample will be a fluid sample and contact will be achieved by introduction of an appropriate volume of the fluid sample onto the array surface, where introduction can be pipette, deposition, and the like.

## Generation of Labeled Target

Targets may be generated by methods known in the art. mRNA can be labeled and used directly as a target, or converted to a labeled cDNA target. Generally, such methods include the use of oligonucleotide primers. Primers that may be employed include oligo dT , random primers, e.g. random hexamers and gene specific primers.

Of particular interest in the generation of labeled target is the use of a set of a representational number of gene specific primers, as described in U.S. Patent Application No. $08 / 859,998$, the disclosure of which is herein incorporated by reference. As the subject sets comprise a representational number of primers, the total number of different primers in any given set will be only a fraction of the total number of different or distinct RNAs in the sample, where the total number of primers in the set will generally not exceed $80 \%$, usually will not exceed $50 \%$ and more usually will not $20 \%$ of the total number of distinct RNAs, usually the total number of distinct messenger RNAs (mRNAs), in the sample. Any two given RNAs in a sample will be considered distinct or different if they comprise a stretch of at least 100 nucleotides in length in which the sequence similarity is less than $98 \%$, as measured using the FASTA algorithm at default settings. As the sets of gene specific primers comprise only a representational number of primers, with physiological sources comprising from 5,000 to 50,000 distinct RNAs, the number of different gene specific primers in the set of gene specific primers will typically range from about 20 to 10,000 , usually from 50 to 2,000 and more usually from 75 to 1500 .

Each of the gene specific primers of the sets described above will be of sufficient length to specifically hybridize to a distinct nucleic acid member of the sample, e.g. RNA or c DNA, where the length of the gene specific primers will usually be at least 8 nt . more usually at least 20 nt and may be as long as 25 nt or longer, but will usually not exceed 50 nt . The gene specific primers will be sufficiently specific to hybridize to complementary template sequence during the generation of labeled nucleic acids under conditions sufficient for first strand cDNA synthesis, which conditions are known by those of skill in the art. The
number of mismatches between the gene specific primer sequences and their complementary template sequences to which they hybridize during the generation of labeled nucleic acids in the subject methods will generally not exceed 20 number $\%$, usually will not exceed 10 number \% and more usually will not exceed 5 number $\%$.

Generally, the sets of gene specific primers will comprise primers that correspond to at least 20 , usually at least 50 and more usually at least 75 distinct genes as represented by distinct mRNAs in the sample, where the term "distinct" when used to describe genes is as defined above, where any two genes are considered distinct if they comprise a stretch of at least 100 nt in their RNA coding regions in which the sequence similarity does not exceed $98 \%$, as determined using the FASTA algorithm at default settings.

The gene specific oligonucleotide primers may be synthesized by conventional oligonucleotide chemistry methods, where the nucleotide units may be: (a) solely nucleotides comprising the heterocyclic nitrogenous bases found in naturally occurring DNA and RNA, e.g. adenine, cytosine, guanine, thymine and uracil; (b) solely nucleotide analogs which are capable of base pairing under hybridization conditions in the course of DNA synthesis such that they function as the above nucleotides found in naturally occurring DNA and RNA, where illustrative nucleotide analogs include inosine, xanthine, hypoxanthine, 1,2diaminopurine and the like; or (c) from combinations of the nucleotides of (a) and nucleotide analogs of (b), where with primers comprising a combination of nucleotides and analogues thereof, the number of nucleotide analogues in the primers will typically be less than 25 and more typically less than 5 . The gene specific primers may comprise reporter or hapten groups, usually 1 to 2 , which serve to improve hybridization properties and simplify detection procedure.

Depending on the particular point at which the gene specific primers are employed in the generation of the labeled nucleic acids, e.g. during first strand cDNA synthesis or following one or more distinct amplification steps, each gene specific primer may correspond to a particular RNA by being complementary or similar, where similar usually means identical, to the particular RNA. For example, where the gene specific primers are employed in the synthesis of first strand cDNA, the gene specific primers will be complementary to regions of the RNAs to which they correspond.

Each gene specific primer can be complementary to a sequence of nucleotides which is unique in the population of nucleic acids, e.g. mRNAs, with which the primers are
contacted. or one or more of the gene specific primers in the set may be complementary to several nucleic acids in a given population, e.g. multiple mRNAs. such that the gene specific primer generates labeled nucleic acid when one or more of set of related nucleic acid species, e.g. species having a conserved region to which the primer corresponds, are present in the sample. Examples of such related nucleic acid species include those comprising: repetitive sequences, such as Alu repeats, A1 repeats and the like; homologous sequences in related members of a gene-family; polyadenylation signals; splicing signals; or arbitrary but conversed sequences.

Depending on the particular nature of the labeled nucleic acid generation step of the subject methods, the gene specific primers may be modified in a variety of ways. One way the gene specific primers may be modified is to include an anchor sequence of nucleotides, where the anchor is usually located $5^{\prime}$ of the gene specific portion of the primer and ranges in length from 10 to 50 nt in length, usually 15 to 40 nt in length. The anchor sequence may comprise a sequence of bases which serves a variety of functions, such as a sequence of bases which correspond to the sequence found in promoters for bacteriophage RNA polymerase, e.g. T7 polymerase, T3 polymerase, SP6 polymerase, and the like; arbitrary sequences which can serve as subsequent primer binding sites; and the like.

Turning now to the methods employing the above sets of gene specific primers, the first step in the subject methods is to obtain a sample of nucleic acids, usually RNAs, from a physiological source, usually a plurality of physiological sources, where the term plurality is used to refer to 2 or more distinct physiological sources. The physiological source of RNAs will typically be eukaryotic, with physiological sources of interest including sources derived single celled organisms such as yeast and multicellular organisms, including plants and animals, particularly mammals, where the physiological sources from multicellular organisms may be derived from particular organs or tissues of the multicellular organism, or from isolated cells derived therefrom. Thus, the physiological sources may be different cells from different organisms of the same species, e.g. cells derived from different humans, or cells derived from the same human (or identical twins) such that the cells share a common genome, where such cells will usually be from different tissue types, including normal and diseased tissue types, e.g. neoplastic, cell types. In obtaining the sample of RNAs to be analyzed from the physiological source from which it is derived, the physiological source may be subjected to a number of different processing steps, where such processing steps
might include tissue homogenation, nucleic acid extraction and the like, where such processing steps are known to the those of skill in the art. Methods of isolating RNA from cells, tissues, organs or whole organisms are known to those of skill in the art and are described in Maniatis et al., Molecular Cloning: A Laboratory Manual (Cold Spring Harbor Press)(1989).

The next step in the subject methods is the generation of labeled nucleic acids representative of the nucleic acid, usually RNA, profile of the physiological source. As mentioned above, a set of gene specific primers is used to generate the labeled nucleic acids from the sample of RNAs, where the labeled nucleic acids generated in this step may serve as "target" in subsequent assays in which the differences in the RNA profiles of at least two sources are analyzed. As used herein, the term "target" refers to single stranded RNA, single stranded DNA and double stranded DNA, where the target is generally greater than 50 nt in length.

The set of primers may be used either in first strand cDNA synthesis or following one or more amplification steps. Furthermore, the actual synthesis of the labeled nucleic acids may be at the same step during which the sets of gene specific primers are employed, or the synthesis of the labeled nucleic acids may be one more steps subsequent to the step in which the sets of gene specific primers are employed.

In a first embodiment of the invention, the set of gene specific primers is used to generate labeled first strand cDNA, where the labeled first strand cDNA is representative of the RNA profile of the physiological source being assayed. The labeled first strand cDNA is prepared by contacting the RNA sample with the primer set and requisite reagents under conditions sufficient for reverse transcription of the RNA template in the sample. Requisite reagents contacted with the primers and RNAs are known to those of skill in the art and will generally include at least an enzyme having reverse transcriptase activity and dNTPs in an appropriate buffer medium.

A variety of enzymes, usually DNA polymerases, possessing reverse transcriptase activity can be used for the first strand cDNA synthesis step. Examples of suitable DNA polymerases include the DNA polymerases derived from organisms selected from the group consisting of a thermophilic bacteria and archaebacteria, retroviruses, yeasts, Neurosporas, Drosophilas, primates and rodents. Preferably, the DNA polymerase will be selected from the group consisting of Moloney murine leukemia virus (M-MLV) as described in United

States Patent No. 4,943,531 and M-MLV reverse transciptase lacking RNaseH activity as described in United States Patent No. 5,405,776 (the disclosures of which patents are herein incorporated by reference), human T-cell leukemia virus type I ( HTLV-I ), bovine leukemia virus ( BLV ), Rous sarcoma virus (RSV ), human immunodeficiency virus (HIV ) and Thermus aquaticus ( Taq ) or Thermus thermophilus (Tth) as described in United States Patent No. $5,322,770$, the disclosure of which is herein incorporated by reference. Suitable DNA polymerases possessing reverse transcriptase activity may be isolated from an organism, obtained commercially or obtained from cells which express high levels of cloned genes encoding the polymerases by methods known to those of skill in the art, where the particular manner of obtaining the polymerase will be chosen based primarily on factors such as convenience, cost, availability and the like.

The various dNTPs and buffer medium necessary for first strand cDNA synthesis through reverse transcription of the primed RNAs may be purchased commercially from various sources, where such sources include Clontech, Sigma, Life Technologies, Amersham, Boehringer-Mannheim. Buffer mediums suitable for first strand synthesis will usually comprise buffering agents, usually in a concentration ranging from 10 to $100 \mu \mathrm{M}$ which typically support a pH in the range 6 to 9 , such as Tris- HCl , HEPES-KOH, etc.; salts containing monovalent ions, such as $\mathrm{KCl}, \mathrm{NaCl}$, etc., at concentrations ranging from 0-200 mM ; salts containing divalent cations like $\mathrm{MgCl}_{2}, \mathrm{Mg}(\mathrm{OAc})$ etc, at concentrations usually ranging from 1 to 10 mM ; and additional reagents such as reducing agents, e.g. DDT, detergents, albumin and the like. The conditions of the reagent mixture will be selected to promote efficient first strand synthesis. Typically the set of primers will first be combined with the RNA sample at an elevated temperature, usually ranging from 50 to $95^{\circ} \mathrm{C}$, followed by a reduction in temperature to a range between about 0 to $60^{\circ} \mathrm{C}$, to ensure specific annealing of the primers to their corresponding RNAs in the sample. Following this annealing step, the primed RNAs are then combined with dNTPs and reverse transcriptase under conditions sufficient to promote reverse transcription and first strand cDNA synthesis of the primed RNAs. By using appropriate types of reagents, all of the reagents can be combined at once if the activity of the polymerase can be postponed or timed to start after annealing of the primer to the RNA.

In this embodiment. one of either the gene specific primers or dNTPs. preferably the dNTPs, will be labeled such that the synthesized cDNAs are labeled. By labeled is meant
that the entities comprise a member of a signal producing system and are thus detectable, either directly or through combined action with one or more additional members of a signal producing system. Examples of directly detectable labels include isotopic and fluorescent moieties incorporated into, usually covalently bonded to, a nucleotide monomeric unit. e.g. dNTP or monomeric unit of the primer. Isotopic moieties or labels of interest include ${ }^{32} P$, ${ }^{33} \mathrm{P},{ }^{35} \mathrm{~S},{ }^{125} \mathrm{I}$, and the like. Fluorescent moieties or labels of interest include coumarin and its derivatives, e.g. 7-amino-4-methylcoumarin, aminocoumarin, bodipy dyes, such as Bodipy FL, cascade blue, fluorescein and its derivatives, e.g. fluorescein isothiocyanate, Oregon green, rhodamine dyes, e.g. texas red, tetramethylrhodamine, eosins and erythrosins, cyanine dyes, e.g. Cy 3 and Cy 5 , macrocyclic chelates of lanthanide ions, e.g. quantum dye ${ }^{\mathrm{TM}}$, fluorescent energy transfer dyes, such as thiazole orange-ethidium heterodimer, TOTAB, etc. Labels may also be members of a signal producing system that act in concert with one or more additional members of the same system to provide a detectable signal. Illustrative of such labels are members of a specific binding pair, such as ligands, e.g. biotin, fluorescein, digoxigenin, antigen, polyvalent cations, chelator groups and the like, where the members specifically bind to additional members of the signal producing system, where the additional members provide a detectable signal either directly or indirectly, e.g. antibody conjugated to a fluorescent moiety or an enzymatic moiety capable of converting a substrate to a chromogenic product, e.g. alkaline phosphatase conjugate antibody; and the like.

In one preferred embodiment, the member of the signal producing system bound to the nucleotide is functional group capable of covalently binding to additional members of the signal producing system to generate a detectable label. Examples of such functional groups or moieties include amino, sulfhydryl, azido, isothiocyanate, sulfoxyl, and the like. The labeled target generated using such nucleotides will thus include one or more, usually a plurality of, functional moieties. For detection, the functional moieties of the modified nucleotides can be labeled by conjugation of a label to the functional moiety. A variety of suitable labels and methods for their conjugation to functional moieties are known to those of skill in the art. Examples include labeling of amino-modified cDNA by a succinimidyl ester of an appropriate dye, e.g. Alexa, Bodipy, or Cy dyes. Alternatively, label can be entrapped or bonded into structures of microscopic-sized particles. These particles can then be conjugated with the functional moieties of the target.

For each sample of RNA, one can generate labeled oligos with the same labels. Alternatively, one can use different labels for each physiological source, which provides for additional assay configuration possibilities, as described in greater detail below.

In a variation of the above embodiment. where desired one can generate labeled RNA instead of labeled first strand cDNA. In this embodiment. first strand cDNA synthesis is carried out in the presence of unlabeled dNTPs and unlabeled gene specific primers. However, the primers are optionally modified to comprise a promotor for an RNA polymerase, such as T7 RNA polymerase, T3 RNA polymerase, SP6 RNA polymerase, and the like. In this embodiment, following first strand cDNA synthesis, the resultant single stranded cDNA is then converted to double stranded cDNA, where the resultant double stranded cDNA comprises the anchor sequence comprising the promoter region. Conversion of the mRNA:cDNA hybrid following first strand synthesis can be carried out as described in Okayama \& Berg, Mol. Cell. Biol. (1982) 2:161-170, and Gubler \& Hoffman, Gene (1983) 25: 253-269, where briefly the RNA is digested with a ribonuclease, such as E.coli RNase $H$, followed by repair synthesis using a DNA polymerase like DNA polymerase I, etc., and E.coli DNA ligase. One may also employ the modification of this basic method described in Wu, R, ed., Methods in Enzymology (1987), vol. 153 (Academic Press). Next, the double stranded cDNA is contacted with RNA polymerase and dNTPs, including labeled dNTPs as described above, to produce linearly amplified labeled ribonucleic acids. For cDNA lacking the anchor sequence comprising a promoter region, a polymerase that does not need a promoter region but instead can initiate RNA strand synthesis randomly from cDNA, such as core fragment of E.Coli RNA polymerase, may be employed.

In another embodiment of the subject invention, the labeled nucleic acid generation step comprises one or more enzymatic amplification steps in which multiple DNA copies of the initial RNAs present in the sample are produced, from which multiple copies of the initial RNA or multiple copies of antisense RNA (aRNA) may be produced, using the polymerase chain reaction, as described in U.S. Pat. No. 4.683,195, the disclosure of which is herein incorporated by reference, in which repeated cycles of double stranded DNA denaturation, oligonucleotide primer annealing and DNA polymerase primer extension are performed, where the PCR conditions may be modified as described in U.S. Pat No. $5,436,149$, the disclosure of which is herein incorporated by reference.

In one embodiment involving enzymatic amplification, the set of gene-specific primers are employed in the generation of the first strand cDNA, followed by amplification of the first strand cDNA to produce amplified numbers of labeled cDNA. In this embodiment, as a set of gene-specific primers is employed in the first strand synthesis step, only a representative proportion of the total RNA in the sample is amplified during the subsequent amplification steps.

Amplification of the first strand cDNA can be conveniently achieved by using a CAPswitch ${ }^{\text {TM }}$ oligonucleotide as described in U.S. Patent Application Serial No. 08/582,562, the disclosure of which is herein incorporated by reference. Briefly, the CAPswitch ${ }^{\text {TM }}$ technology uses a unique CAPswitch ${ }^{\text {™ }}$ oligonucleotide in the first strand cDNA synthesis followed by PCR amplification in the second step to generate a high yield of ds cDNA. When included in the first-strand cDNA synthesis reaction mixture, the CAPswitch ${ }^{\text {TM }}$ oligonucleotide serves as a short extended template. When reverse transcriptase stops at the 5 ' end of the mRNA template in the course of first strand cDNA synthesis it switches templates and continues DNA synthesis to the end of the CAPswitch ${ }^{\text {TM }}$ oligonucleotide. The resulting ss cDNA incorporates at the $3^{\prime}$ end, sequence which is complimentary to complete $5^{\prime}$ end of the mRNA and the CAPswitch ${ }^{\text {TM }}$ oligonucleotide sequence.

Of particular interest as the CAPswitch ${ }^{\text {M }}$ oligonucleotide are oligonucleotides having the following formula:

$$
5^{\prime}-\mathrm{dN} 1-\mathrm{dN} 2-. . . \mathrm{dNm}-\mathrm{rN} 1-\mathrm{rN} 2 \ldots . . \mathrm{rNn}-3{ }^{\prime}
$$

wherein:
dN represents a deoxyribonucleotide selected from among dAMP, dCMP, dGMP and dTMP;
m represents an integer 0 and above, preferably from 10 to 50 ;
rN represents a ribonucleotide selected from the group consisting of AMP, CMP, GMP and UMP, preferably GMP; and
n represents an integer 0 and above, preferably from 3 to 7.

The structure of the CAPswitch ${ }^{\text {TM }}$ oligonucleotide may be modified in a number of ways. such as by replacement of 1 to 10 nucleotides with nucleotide analogs, incorporation
of terminator nucleotides, such as 3 '-amino NMP. $3^{\prime}$-phosphate NMP and the like. or non-natural nucleotides which can improve efficiency of the template switching reaction but still retain the main function of the CAPswitch ${ }^{\text {TM }}$ oligonucleotide i.c. CAP-depended extension of full-length cDNA by reverse transcriptase using CAPswitch ${ }^{\text {TM }}$ oligonucleotide as a template.

In using the CAPswitch ${ }^{\text {TM }}$ oligonucleotide, first strand cDNA synthesis is carried out in the presence of a set of gene specific primers and a CAPswitch ${ }^{\top \mathrm{M}}$ oligonucleotide, where the gene specific primers have been modified to comprise an arbitrary anchor sequence at their 5 ' ends. The first strand cDNA is then combined with primer sequences complementary to: (a) all or a portion of the CAPswitch ${ }^{\text {TM }}$ oligonucleotide and (b) the arbitrary anchor sequence of the gene specific primers and additional PCR reagents, such as dNTPs, DNA polymerase, and the like, under conditions sufficient to amplify the first strand cDNA. Conveniently, PCR is carried out in the presence of labeled dNTPs such that the resultant, amplified cDNA is labeled and serves as the labeled or target nucleic acid. Labeled nucleic acid can also be produced by carrying out PCR in the presence of labeled primers, where either or both the CAPswitch ${ }^{\text {TM }}$ oligonucleotide complementary primer and anchor sequence complementary primer may be labeled. In yet an alternative embodiment, instead of producing labeled amplified cDNA, one may generate labeled RNA from the amplified ds cDNA, e.g. by using an RNA polymerase such as E.coli RNA polymerase, or other RNA polymerases requiring promoter sequences, where such sequences may be incorporated into the arbitrary anchor sequence.

Instead of using the set of gene specific primers in the first strand cDNA synthesis step followed by subsequent amplification of only a representative fraction of the total number of distinct RNA species in the sample, one may also amplify all of the RNAs in the sample and use the set of gene specific primers to generate labeled nucleic acid following amplification. This embodiment may find use in situations where the RNA of interest to be amplified is known or postulated to be in small amounts in the sample.

In this embodiment, first strand synthesis is carried out using: (a) an oligo dT primer that usually comprises an arbitrary anchor sequence at its 5 ' end and (b) a CAPswitch ${ }^{\text {TM }}$ oligonucleotide. During first strand synthesis the oligo(dT) anneals to the polyA tail of the mRNA in the sample and synthesis extends beyond the 3 ' end of the RNA to include the CAPswitch ${ }^{\text {™ }}$ oligonucleotide, yielding a first strand cDNA comprising an arbitrary
sequence at its $5^{\prime}$ end and a region complementary to the CAPswitch ${ }^{T M}$ oligonucleotide at its 3 ' end. The length of the dT primer will typically range from 15 to 30 nts , while the arbitrary anchor sequence or portion of the primer will typically range from 15 to 25 nt in length.

Following first strand synthesis, the cDNA is amplified by combining the first strand cDNA with primers that correspond at least partially to the anchor sequence and the CAPswitch ${ }^{\text {TM }}$ oligonucleotide under conditions sufficient to produce an amplified amount of the cDNA. Labeled nucleic acid is then produced by contacting the resultant amplified cDNA with a set of gene specific primers, a polymerase and dNTPs, where at least one of the gene specific primers and dNTPs are labeled.

When employed to generate target, as described above, the gene specific primers of the sets of primers according to the subject invention are typically chosen according to a number of different criteria. In some embodiments of the invention, primers of interest for inclusion in the set include primers corresponding to genes which are typically differentially expressed in different cell types, in disease states, in response to the influence of external agents, factors or infectious agents, and the like. In other embodiments, primers of interest are primers corresponding to genes which are expected to be, or already identified as being, differentially expressed in different cell, tissue or organism types. Preferably, at least 2 different gene functional classes will be represented in the sets of gene specific primers, where the number of different functional classes of genes represented in the primer sets will generally be at least 3 , and will usually be at least 5 . Gene functional classes of interest include oncogenes; genes encoding tumor suppressors; genes encoding cell cycle regulators; stress response genes; genes encoding ion channel proteins; genes encoding transport proteins; genes encoding intracellular signal transduction modulator and effector factors; apoptosis related genes; DNA synthesis/recombination/repair genes; genes encoding transcription factors; genes encoding DNA-binding proteins; genes encoding receptors. including receptors for growth factors, chemokines, interleukins, interferons, hormones, neurotransmitters, cell surface antigens, cell adhesion molecules etc.; genes encoding cellcell communication proteins, such as growth factors, cytokines. chemokines, interleukins, interferons, hormones etc.; and the like. Less preferred are gene specific primers that are subject to formation of strong secondary structures with less than $-10 \mathrm{kcal} / \mathrm{mol}$; comprise stretches of homopolymeric regions, usually more than 5 identical nucleotides; comprise
more than 3 repetitive sequences; have high. e.g. more than $80 \%$, or low, e.g. less than $30 \%$, GC content etc.

The particular genes represented in the set of gene specific primers will necessarily depend on the nature of physiological source from which the RNAs to be analyzed are derived. For analysis of RNA profiles of eukaryotic physiological sources, the genes to which the gene specific primers correspond will usually be Class II genes which are transcribed into RNAs having 5' caps, e.g. 7-methyl guanosine or 2,2,7-trimethylguanosine, where Class II genes of particular interest are those transcribed into cytoplasmic mRNA comprising a 7 -methyl guanosine 5 ' cap and a polyA tail.

For analysis of RNA profiles of mammalian physiological sources, of particular interest are gene specific primers corresponding to the functional gene classes listed above. For analysis of RNA profiles of human physiological sources, the gene specific primers of particular interest are the gene specific primers identified in Table 1 as SEQ ID NO:01 to SEQ ID NO:1372, of U.S. Application Serial No. 08/859,998, the disclosure of which is herein incorporated by reference, where sets of these primers will usually include at least 20 and more usually at least 50 of these specific sequences.

Particular sets of primers of interest in the subject invention are those sets of primers that include primers capable of amplifying at least a portion of the unique polynucleotides present on the arrays with which the target is to be employed. By at least a portion is meant at least about 10 , usually at least about 20 and more usually at least about 25 number \% (where number is the number of different unique polynucleotides on the array). For examples, sets of primers that include primers capable of amplifying at least a portion of the unique polynucleotides listed in Table 1, supra, are of interest. Similarly sets of primers capable of amplifying at least a portion of the unique polynucleotides listed in Tables 2 to 8, supra, are also of interest.

In a particularly preferred embodiment, the gene specific primers are preferably those primers that correspond to the different polynucleotide spots on the array that is used in the hybridization assay. Thus, one will preferably employ gene specific primers for each different polynucleotide that is present on the array, so that if the gene is expressed in the particular cell or tissue being analyzed, labeled target will be generated from the sample for that gene. In many embodiments in which the subject arrays are employed, the gene specific primers used to generate the target from the human cell or tissue being analyzed will have
the same sequence as the gene specific primers used to generate the polynucleotide probes present on the array. In this manner, if a particular gene present on the array is expressed in a particular sample, the appropriate target will be generated and subsequently identified. Representative sets of primers falling within this particularly preferred embodiment include:

SET

1

2

3

DESCRIPTION
I pair of primers capable of amplifying each polynucleotide listed in Table I, supra, as well one set of primers capable of amplifying each of the complementary sequences of each of the polynucleotides listed in Table I.

I pair of primers capable of amplifying each polynucleotide listed in Table 2, supra, as well one set of primers capable of amplifying each of the complementary sequences of each of the polynucleotides listed in Table 2.

I pair of primers capable of amplifying each polynucleotide listed in Table 3, supra, as well one set of primers capable of amplifying each of the complementary sequences of each of the polynucleotides listed in Table 3.

## Hybridization and Detection

As mentioned above, following preparation of the target nucleic acid from the tissue or cell of interest, the target nucleic acid is then contacted with the array under hybridization conditions, where such conditions can be adjusted, as desired, to provide for an optimum level of specificity in view of the particular assay being performed. Suitable hybridization conditions are well known to those of skill in the art and reviewed in Maniatis et al, supra and WO 95/21944. In analyzing the differences in the population of labeled target nucleic acids generated from two or more physiological sources using the arrays described above, each population of labeled target nucleic acids are separately contacted to identical probe arrays or together to the same array under conditions of hybridization, preferably under stringent hybridization conditions (for example, at $50^{\circ} \mathrm{C}$ or higher and 0.1 XSSC $(15 \mathrm{mM}$ sodium chloride $/ 01.5 \mathrm{mM}$ sodium citrate)), such that labeled target nucleic acids hybridize to complementary probes on the substrate surface.

Where all of the target sequences comprise the same label, different arrays will be employed for each physiological source (where different could include using the same array at different times). Alternatively, where the labels of the targets are different and
distinguishable for each of the different physiological sources being assayed, the opportunity arises to use the same array at the same time for each of the different target populations. Examples of distinguishable labels are well known in the art and include: two or more different emission wavelength fluorescent dyes, like Cy 3 and Cy 5 , two or more isotopes with different energy of emission, like ${ }^{32} \mathrm{P}$ and ${ }^{33} \mathrm{P}$, labels which generate signals under different treatment conditions, like temperature, pH , treatment by additional chemical agents, etc., or generate signals at different time points after treatment. Using one or more enzymes for signal generation allows for the use of an even greater variety of distinguishable labels, based on different substrate specificity of enzymes (alkaline phosphatase/peroxidase).

Following hybridization, non-hybridized labeled nucleic acid is removed from the support surface, conveniently by washing, generating a pattern of hybridized nucleic acid on the substrate surface. A variety of wash solutions are known to those of skill in the art and may be used.

The resultant hybridization patterns of labeled nucleic acids may be visualized or detected in a variety of ways, with the particular manner of detection being chosen based on the particular label of the target nucleic acid, where representative detection means include scintillation counting, autoradiography, fluorescence measurement, colorimetric measurement, light emission measurement and the like.

Following detection or visualization, the hybridization patterns may be compared to identify differences between the patterns. Where arrays in which each of the different probes corresponds to a known gene are employed, any discrepancies can be related to a differential expression of a particular gene in the physiological sources being compared.

## Utility

The subject methods find use in, among other applications, differential gene expression assays. Thus, one may use the subject methods in the differential expression analysis of: (a) diseased and normal tissue, e.g. neoplastic and normal tissue, (b) different tissue or tissue types; (c) developmental stage; (d) response to external or internal stimulus; (e) response to treatment; and the like. The subject arrays therefore find use in broad scale expression screening for drug discovery and research, such as the effect of a particular active agent on the expression pattern of genes in a particular cell, where such information can be
used to reveal drug toxicity, carcinogenicity, etc., environmental monitoring, disease research and the like.

## Kits

Also provided are kits for performing analyte binding assays using the subject devices, where kits for carrying out differential gene expression analysis assays are preferred. Such kits according to the subject invention will at least comprise the subject arrays. The kits may further comprise one or more additional reagents employed in the various methods, such as primers for generating target nucleic acids, including a set of gene specific primers according to the subject invention, e.g. primer sets 1 to 9 described above, dNTPs and/or rNTPs, which may be either premixed or separate, one or more uniquely labeled dNTPs and/or rNTPs, such as biotinylated or Cy3 or Cy5 tagged dNTPs, or other post synthesis labeling reagent, such as chemically active derivatives of fluorescent dyes, enzymes, such as reverse transcriptases, DNA polymerases, and the like, various buffer mediums, e.g. hybridization and washing buffers, prefabricated probe arrays, labeled probe purification reagents and components, like spin columns, etc., signal generation and detection reagents, e.g. streptavidin-alkaline phosphatase conjugate, chemifluorescent or chemiluminescent substrate, and the like.

The following examples are offered by way of illustration and not by way of limitation

## EXPERIMENTAL

## Example 1-Generation of human cDNA array

686 cDNA fragments corresponding 686 different human genes were amplified from quick-clone cDNA (CLONTECH) in 686 separate test tubes using a combination of sense and antisense gene-specific primers: (Set No. 9, described supra). Amplification was conducted in a $100-\mu \mathrm{l}$ volume containing $2 \mu \mathrm{l}$ of mixture of 10 Quick-clone cDNA from placenta, brain, liver, lung, leukocytes, spleen, skeletal muscle, testis, kidney and ovary (CLONTECH), 40 mM Tricine-KOH (pH 9.2 at $22^{\circ} \mathrm{C}$ ), $3.5 \mathrm{mM} \mathrm{Mg}(\mathrm{OAc})_{2}, 10 \mathrm{mM}$ KOAc.
$75 \mu \mathrm{~g} / \mathrm{ml}$ BSA, $200 \mu \mathrm{M}$ of each dATP, dGTP, dCTP and dTTP, $0.2 \mu \mathrm{M}$ of each sense and antisense gene-specific primers and $2 \mu$ l of KlenTaq Polymerase mix. Temperature parameters of the PCR reactions were as follows: 1 min at $95^{\circ} \mathrm{C}$ followed by $20-35$ cycles of $95^{\circ} \mathrm{C}$ for 15 sec and $68^{\circ} \mathrm{C}$ for 2 min ; followed by a $10-\mathrm{min}$ final extension at $68^{\circ} \mathrm{C} . \mathrm{PCR}$ products were examined on $1.2 \%$ agarose $/ E t B r$ gels in $1 \times$ TBE buffer. As a DNA size marker a 1 Kb DNA Ladder was used. ds cDNA was then precipitated by addition of a half volume of 4 M ammonium acetate (about $35 \mu \mathrm{l}$ ) and 3.7 volumes of $95 \%$ ethanol (about 260 $\mu \mathrm{l}$ ). After vortexing, the tube was immediately centrifuged at 14,000 r.p.m. in a microcentrifuge for 20 min . The pellet was washed with $80 \%$ ethanol without vortexing, centrifuged as above for 10 min , air dried, and dissolved in $10 \mu \mathrm{l}$ of deionized water. Yield of ds cDNA after the amplification step was about $5 \mu \mathrm{~g}$. The ds cDNA fragments for all 686 genes were cloned into TA-cloning vector using the manufacturer's recommendations (Invitrogen) and identity of the clones was confirmed by sequence analysis. The ds cDNA inserts with the sequence corresponding 686 genes were amplified by PCR using a combination of antisense and sense gene-specific primers, as described above. The ds cDNA was denatured by adding $1 \mu \mathrm{l}$ of 10 X denaturing solution ( $1 \mathrm{M} \mathrm{NaOH}, 10 \mathrm{mM}$ EDTA) and incubating at $65^{\circ} \mathrm{C}$ for 20 min . All cDNA probes were transferred in 384 -well plate and loaded on positively charged nylon membrane (Schleher \& Schull) using 384 pin tool and Biomek 2000 (Beckman) robot. The resultant array is described in Table 1.

## Example 2- Generation of ${ }^{32}$ P-labeled oligonucleotides during first strand cDNA

 synthesisStep A. cDNA synthesis/Labeling Procedure
$1 \mu \mathrm{~g}$ of polyA+RNA or total RNA was converted into ${ }^{32}$ - - labeled first-strand cDNA as follows. A sufficient volume of master mix for all labeling reactions and 1 extra reaction was prepared as follows to ensure sufficient volume. For each $10-\mu$ l labeling reaction, the following reagents were mixed:

[^0]$8 \mu \mathrm{l} \quad$ Final volume
Next, the following reagents were combined in a $0.5-\mathrm{ml}$ PCR test tube:
$1 \mu \mathrm{~g}(1-2 \mu \mathrm{l}) \quad$ polyA + RNA sample
$|\mu| \quad 10 x$ gene-specific primers mix ( $0.2 \mu \mathrm{M}$ of each oligonucleotide ID No.
$2,4,6,8,10,12, \ldots . .1372$ from Table I of U.S. Patent Application Serial No.
$08 / 859,998$, the discosure of which is herein incorporated by reference.)

As a control, in separate test tube were mixed $1 \mu \mathrm{~g}$ of polyA+RNA sample with $1 \mu \mathrm{l}$ of oligo dT primer (CDS1, $5^{\prime}-\mathrm{d}\left(\right.$ TCTAGAATTCAGCGGCCGC(T) $\left.{ }_{30} \mathrm{VN}\right) \cdot 3^{\prime}$
(where $\mathrm{V}=\mathrm{G}$ or A or $\mathrm{C} ; \mathrm{N}=\mathrm{G}$ or A or T or C )

For each tube, $\mathrm{ddH}_{2} 0$ was added to a final volume of $3 \mu \mathrm{l}$ and the contents were mixed and spun briefly in a microcentrifuge. The tubes were then incubated in a preheated PCR thermocycler at $70^{\circ} \mathrm{C}$ for 2 min . The temperature in thermocycle was reduced down to $50^{\circ} \mathrm{C}$ and the tube contents were incubated for $2 \mathrm{~min} .8 \mu \mathrm{l}$ of master mix as prepared above were added to each reaction test tube. The contents of the test tubes were then mixed by gentle pipetting. The tubes were then incubated in a PCR thermocycler for 20 min at $50^{\circ} \mathrm{C}$. The reaction was then stopped by adding $1 \mu \mathrm{l}$ of 10 X termination mix ( 0.1 M EDTA, 1 $\mathrm{mg} / \mathrm{ml}$ glycogen).

## Step B. Column Chromatography

The ${ }^{32} \mathrm{P}$-labeled cDNAs were separated from unincorporated ${ }^{32} \mathrm{P}$-labeled nucleotides and small ( $<0.1-\mathrm{kb}$ ) cDNA fragments using the following procedure for each test tube. A CHROMA SPIN-200 column (CLONTECH, Palo Alto. CA) was placed into a $1.5-\mathrm{ml}$ microcentrifuge tube, the water was allowed to drain through the column by gravity flow until the surface of the gel beads emerged in the column matrix. The sample was then applied to the center of the gel bed's flat surface and allowed to be fully absorbed into the resin bed. $25 \mu \mathrm{l}$ of $\mathrm{ddH}_{2} \mathrm{O}$ were then applied and allowed to completely drain out of the column. $200 \mu \mathrm{l}$ of $\mathrm{ddH}_{2} \mathrm{O}$ were then applied and allowed to completely drain out of the column until there was no liquid left above the resin bed. The column was then transferred to a clean $1.5-\mathrm{ml}$ microcentrifuge tube.

To collect the first fraction, $100 \mu \mathrm{l}$ of $\mathrm{ddH}_{2} \mathrm{O}$ were added to the column and allowed to completely drain out of the column. The second, third and fourth fractions were collected in analogous fashion. The tubes with fractions 1-4 were then placed in scintillation counter empty vials, and Cherenkov counts for each fraction were obtained in the tritium channel. The fractions which showed the highest Cerenkov counts were pooled.

## Example 3- Generation of Cy 3 -labeled hybridization polynucleotide target from polyA+RNA using postsynthesis labelling procedure

In this procedure for generating labeled cDNA target, polyA+RNA is first converted into cDNA that has primary amino groups which are subsequently coupled with Cy 3 succinimide ester. This technology allows for a significant increase (about 10 fold) in activity of labeled polynucleotide target and therefore increases the overall sensitivity of detection of gene expression. The same procedure can be used for labeling two (or more) samples of RNA. In this case the cDNA synthesis step was the same for both samples but at the labeling step, each cDNA sample was labeled by different and distinguishable labels, e.g. Cy 3 and Cy5, Alexa 532 and Bodipy TR, Fluorescein and tetramethyl rhodamine, etc. Each labeled probe was purified separately by column chromatography and, after normalization, were combined together in equal ratio and hybridized with a cDNA array. After hybridization, the detection procedure revealed both dye-labeled hybridized target simultaneously, based on the different spectral characteristics (emission wavelength) of the fluorescent labels.

## A. cDNA synthesis

The $10-\mu 1$ reaction described below converted $1 \mu \mathrm{~g}$ of polyA+RNA into aminomodified first-strand cDNA.

For each cDNA synthesis reaction:

1. Enough master mix for all labeling reactions and 1 extra reaction was prepared to ensure sufficient volume.

For each $10-\mu$ l labeling reaction, the following reagents were mixed:

$$
\begin{array}{ll}
2 \mu \mathrm{l} & 5 \mathrm{X} \text { First-strand buffer ( } 250 \mu \mathrm{M} \text { Tris-HCI pH8.3: } 375 \mathrm{mM} \mathrm{KCl}: 15 \mathrm{mM} \mathrm{MgCl} \text { ) }) \\
1 \mu \mathrm{l} & 10 \mathrm{XdNTP} \text { mix ( } 500 \mu \mathrm{M} \mathrm{dGTP}, 500 \mu \mathrm{M} \mathrm{dCTP}, 500 \mu \mathrm{MdATP}, 100 \mu \mathrm{M} \mathrm{dTTP} .
\end{array}
$$

and $100 \mu \mathrm{M}$ allylamino dUTP )
$|\mu| \quad\left[\alpha-{ }^{-12} \mathrm{P}\right] \mathrm{dATP}$ (Amersham, $3000 \mathrm{Ci} / \mathrm{mmol}, 10 \mathrm{mCi} / \mathrm{ml}$ )
$3 \mu \mathrm{l} \quad \mathrm{H}_{2} \mathrm{O}$
$1 \mu \mathrm{l}$ MMLV reverse transcriptase (Amersham. 200 units/ul)
$8 \mu$ I Final volume
2. The following was combined in a $0.5-\mathrm{ml}$ PCR test tube:

```
l \mug(1-2 \mul) polyA+RNA sample
| | 10x gene-specific primers mix ( 0.2 uM of each oligonucleotide ID No.
    2,4,6,8,10,12,\ldots..... 1372) (from Table I of U.S. Patent Application No.
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$08 / 859.998$, the disclosure of which is herein incorporated by reference.)

As a control in separate test tube $1 \mu \mathrm{~g}$ of polyA+RNA sample was mixed with $1 \mu \mathrm{l}$ of oligo dT primer (SEQ ID NO. 1373 from Table 1 of U.S. Application No. 08/859,998).
3. $\mathrm{ddH}_{2} \mathrm{O}$ was added to a final volume of $3 \mu \mathrm{l}$.
4. The contents were mixed and the tubes were spun briefly in a microcentrifuge.
5. The tubes were incubated in preheated PCR thermocycler at $70^{\circ} \mathrm{C}$ for 2 min .
6. The temperature in the thermocycle was reduced down to $50^{\circ} \mathrm{C}$ and incubate for 2 min.
7. $8 \mu \mathrm{l}$ of master mix were added to each reaction test tube.
8. The contents of the test tubes were mixed by gentle pipeting.
9. The tubes were incubated in a PCR thermocycler for 30 min at $50^{\circ} \mathrm{C}$.
10. The reaction was stopped by increasing temperature up to $70^{\circ} \mathrm{C}$ for 5 min , then cooled to $37^{\circ} \mathrm{C}$.
11. $1 \mu \mathrm{l}$ of $\mathrm{RNase} \mathrm{H}(10$ units $/ \mu \mathrm{l})$ was added and the tubes were incubated at $37^{\circ} \mathrm{C}$ for 15 min.
12. The reaction was stopped by adding $40 \mu \mathrm{l}$ of termination $\operatorname{mix}$ ( 0.3 M sodium acetate. pH 5.0, 1 mMEDTA).
13. An equal volume ( $50 \mu \mathrm{I}$ ) of phenol/chlorophorm/isoamyl alcohol mix ( $1: 1: 1 / 24 \mathrm{v} / \mathrm{v}$ ) was added and extraction was performed. Phases were separated by centrifugation at 14.000 rpm for 10 min .
14. Upper water phase was collected and cDNA was precipitated by adding 2.5 volumes (about $120 \mu \mathrm{l}$ ) of ethanol.
15. The precipitate was collected by centrifugation at $14,000 \mathrm{rpm}$ for 10 min , the supernatant removed and the precipitate was washed with $80 \%$ ethanol.
16. The precipitate was air dried and dissolved in $10 \mu \mathrm{l}$ of 0.1 M sodium bicarbonate buffer, pH 9.0 .

Step B. Post synthesis labeling procedure.

1. 1 mg of Cy 3 succinimide ester was dissolved in $10 \mu \mathrm{l}$ of dimethyl sulfoxide and 10 $\mu l$ of amino-modified cDNA generated at step 16 was added to it.
2. The mixture was incubated at room temperature overnight.

## Step C. Column Chromatography

To purify the Cy3-labeled cDNAs from the unconjugated label, the following was performed for each test tube:

1. CHROMA SPIN-200 column (CLONTECH) was removed from refrigerator and allowed to warm at room temperature for about 1 hour. The column was inverted several times to completely resuspend the gel matrix. (Note: Check for air bubbles in the column matrix. If bubbles are visible, resuspend the matrix in the in the column buffer ( $\mathrm{ddH}_{2} \mathrm{O}$ ) by inverting the column again).
2. The bottom cap from the column was removed, and then the top cap was slowly removed.
3. The column was placed into a $1.5-\mathrm{ml}$ microcentrifuge tube.
4. The water was allowed to drain through the column by gravity flow until the surfaces of the gel beads in the column matrix were visible. (The top of the column matrix should be at the $0.75-\mathrm{ml}$ mark on the wall of the column. If the column contains much less matrix, adjust the volume of the matrix to 0.75 ml mark using matrix from another column.)
5. The collected water was discarded.
6. The sample was applied to the center of the gel bed's flat surface and allowed to be fully absorbed into the resin bed. Care was taken not allow any sample to flow along the inner wall of the column.
7. $25 \mu \mathrm{l}$ of $\mathrm{ddH}_{2} 0$ were applied and allowed to completely drain out of the column.
8. Apply $200 \mu \mathrm{l}$ of $\mathrm{ddH}_{2} \mathrm{O}$ and allow the buffer to completely drain out of the column until there was no liquid left above the resin bed.
9. The column was transfered to a clean $1.5-\mathrm{ml}$ microcentrifuge tube.
10. $100 \mu \mathrm{l}$ of $\mathrm{ddH}_{2} \mathrm{O}$ were added to the column and allowed to completely drain out of the column.
11. The second, third and fourth fractions were collected by repeating steps 9-10.
12. Cherenkov counts were obtained for each fraction by counting the entire sample in the tritium channel.
13. The fractions (usually fractions 2-3) which showed highest Cerenkov counts were pooled. Waste column and the fractions (usually fraction 1 and 4) which showed less than $10 \%$ counts from peak fractions.

## Example 4-Hybridization ${ }^{32}$ P-labeled cDNA Target with cDNA Array

A solution of ExpressHyb ${ }^{\text {TM }}$ (CLONTECH) and sheared salmon testes DNA (Sigma) was prepared by prewarming 15 ml of ExpressHyb ${ }^{\mathrm{TM}}$ at $50-60^{\circ} \mathrm{C}$, heating 1.5 mg of sheared salmon testes DNA at $95-100^{\circ} \mathrm{C}$ for 5 min followed by chilling quickly on ice, and combining the resultant heat-denatured sheared salmon testes DNA with the prewarmed ExpressHyb ${ }^{\mathrm{TM}}$.

A cDNA Array as produced in Example 1 above was then placed in a hybridization bottle and 10 ml of the solution prepared above was added to the bottle. Prehybridization was performed for 30 min with continuous agitation at $72^{\circ} \mathrm{C}$. Labeled cDNA probe (Example 1, about 200 ul , total about $2-5 \times 10^{6} \mathrm{cpm}$ ) with $1 / 10$ th of the total volume ( about 22 ul ) of 10 x denaturing solution ( $1 \mathrm{M} \mathrm{NaOH}, 10 \mathrm{mM}$ EDTA) was mixed and incubated at $65^{\circ} \mathrm{C}$ for 20 min . $5 \mu \mathrm{l}(1 \mu \mathrm{~g} / \mathrm{ul})$ of human Cot-1 DNA was then added. and an equal volume (about $225 \mu \mathrm{l}$ ) of $2 \times$ Neutralizing solution ( $1 \mathrm{M} \mathrm{NaHPO} 4, \mathrm{pH} 7.0$ ) was added and incubation continued at $65^{\circ} \mathrm{C}$ for 10 min . The mixtures were then combined and thoroughly mixed.

The prehybridization solution was replaced with the solution comprising the labeled oligonucleotide as prepared above and allowed to hybridize overnight with continuous agitation at $65^{\circ} \mathrm{C}$. Following hybridization, the hybridization solution was carefully removed
and discarded, replaced with 200 ml of Wash Solution 1 (2X SSC. 1\% SDS). The array was washed for 20 min with continuous agitation at $65^{\circ} \mathrm{C}$. Washing was repeated four times.

Two additional $20-\mathrm{min}$ washes were then performed in 200 ml of prewarmed Wash Solution $2(0.1 \mathrm{X} \mathrm{SSC}, 0.5 \% \mathrm{SDS})$ with continuous agitation at $65^{\circ} \mathrm{C}$. Using forceps, the cDNA array was removed from the container and excess wash solution was removed by shaking.

The damp membrane was immediately wrapped in plastic wrap, mounted on Whatman paper ( 3 mm Chr ) and exposed to x -ray film at $-70^{\circ} \mathrm{C}$ with an intensifying screen.

## Example 5 -Comparison Between Using Sets of Gene Specific Primers and oligo dT

${ }^{32}$ P-labeled cDNA target were synthesized by M-MLV reverse transcriptase from a mixture 588 antisense gene-specific primers (B) or oligo $\mathrm{dT}(\mathrm{A})$ using placenta polyA+RNA as a template as described in Example 2. Primer extension products generated by reverse transcription were purified by gel filtration as described in Example 2 and hybridized separately with two cDNA arrays comprising 588 human genes under identical conditions as described in Example 4. Signals which can be detected by using cDNA target generated using the set of gene specific primers but can not be detected by using conventional target generated with oligo dT primers were observed. Note, the level of non-specific background detected as signal generated by membrane alone outside of the regions with immobilized probes generated by target generated using oligo dT primers was significantly higher in comparison with the background generated by the target generated by using the sets of gene specific primers.

Example 6-Generation of cDNA array probe immobilized on glass slides.
50 cDNA fragments corresponding to 50 different human genes were amplified from plasmid clones containing corresponding cDNA fragments in 96 well plates using combination of vector primer ID No. 1376 and ID No. 1377 or sense and antisense genespecific primers: ID No. $1+2,3+4,5+6,7+8, \ldots . .100+101$ (from Table 1 of U.S. Patent Application No. 08/859,998, the disclosure of which is herein incorporated by reference). Amplification was conducted in a $400-\mu \mathrm{l}$ volume containing 2 ng of plasmid DNA. 40 mM Tricine- $\mathrm{KOH}\left(\mathrm{pH} 9.2\right.$ at $22^{\circ} \mathrm{C}$ ), $3.5 \mathrm{mM} \mathrm{Mg}(\mathrm{OAc})_{2}, 10 \mathrm{MM}$ KOAc. $75 \mu \mathrm{~g} / \mathrm{ml} \mathrm{BSA}, 200 \mu \mathrm{M}$
of each dATP. dGTP. dCTP and dTTP, $0.2 \mu \mathrm{M}$ of each primers and $2 \mu \mathrm{l}$ of KlenTaq Polymerase mix (CLONTECH). Temperature parameters of the PCR reactions were as follows: 1 min at $95^{\circ} \mathrm{C}$ followed by 30 cycles of $95^{\circ} \mathrm{C}$ for 15 sec and $68^{\circ} \mathrm{C}$ for 2 min ; followed by a $10-\mathrm{min}$ final extension at $68^{\circ} \mathrm{C}$. PCR products were examined on $1.2 \%$ agarose $/ \mathrm{EtBr}$ gels in $1 \times$ TBE buffer. As a DNA size marker, a 1 Kb DNA Ladder was used. ds cDNA was then precipitated by addition of a $10 \%$ volume of 3 M sodium acetate ( $\mathrm{pH} 5-0$ ) (about $40 \mu \mathrm{l}$ ) and 2.5 volumes of $96 \%$ ethanol (about 1 ml ). After vortexing, the tube was immediately centrifuged at 14,000 r.p.m. in a microcentrifuge for 20 min . The pellet was washed with $80 \%$ ethanol without vortexing, centrifuged as above for 10 min , air dried, and dissolved in $10 \mu \mathrm{l}$ of deionized water. Yield of ds cDNA after amplification step was about $20 \mu \mathrm{~g}$. The ds cDNA was solved in $10 \mu \mathrm{l}$ of distilled water, $10 \mu \mathrm{l}$ of 1 M sodium carbonate buffer, pH 9.5 , was added and the ds cDNA was denaturated by heating at $94^{\circ} \mathrm{C}$ for 5 min and cooled down. The treated glass slides were prepared as following: Glass slides were cleaned overnight in $25 \%$ solution of nitric acid at room temperature, washed 3 times by acetone, treated with $1 \%$ aminopropyl-trimethoxysilane for 3 hrs at room temperature, washed two times with acetone, heated at $120^{\circ} \mathrm{C}$ for 6 hrs and then treated with $0.2 \%$ of para-phenylendiisothiocyanate (95:5 acetone-water solution) at room temperature for 3 hrs , then washed two times by acetone and dried in vacuum with desiccant. All cDNA probes were transferred in 384 -well plate and printed on treated glass slides using 384 pin tool and Biomek 2000 (Beckman) robot. After printing, the arrays were incubated in wet chamber at $37^{\circ} \mathrm{C}$ overnight, then ultraviolet-cross linked to the surface by subjecting the slides to 30 mJ of energy (Stratagene Stratalinker). The arrays were washed with $1 \%$ of sodium borohydrate in 0.1 M NaOH , then washed 3 times in distilled water, dried in vacuum and stored with desiccant.

## Example 7- Hybridization Cy 3 -labeled cDNA Target (or Cy3/Cy5 labeled cDNA targets) with glass cDNA array

1. A solution of ExpressHyb (CLONTECH) and sheared salmon testes DNA (Sigma) was prepared as follows:
a. 5 ml of ExpressHyb ${ }^{\text {TM }}$ was prewarmed at $50-60^{\circ} \mathrm{C}$.
b. $\quad 0.5 \mathrm{mg}$ of the sheared salmon testes DNA was heated at $95-100^{\circ} \mathrm{C}$ for 5 $\min$, and then chilled quickly on ice.
c. Heat-denatured sheared salmon testes DNA was mixed with prewarmed ExpressHyb.
2. The glass cDNA array was placed in a hybridization container, and 1 ml of the solution prepared in step 1 above was added.
3. Prehybridization was conducted for 5 min with continuous agitation at $65^{\circ} \mathrm{C}$.
4. Labeled cDNA probe as prepared in example 3, step C13, above, (about $200 \mu \mathrm{l}$ ) was mixed with $2 \mu \mathrm{l}(1 \mu \mathrm{~g} / \mu \mathrm{l})$ of human Cot- I DNA , and denaturated at $99^{\circ} \mathrm{C}$ for 2 min.
5. The mixture prepared in Step 4 was added to the hybridization box from Step 3 and the two solutions were mixed together thoroughly. The container was sealed by sealing tape.
6. Hybridization was allowed to proceed overnight with continuous agitation at $65^{\circ} \mathrm{C}$.
7. The hybridization solution was carefully removed and discarded in an appropriate container, and replaced with 10 ml of Wash Solution 1 (2X SSC, $0.1 \%$ SDS). The array was washed for 10 min with continuous agitation at $65^{\circ} \mathrm{C}$. The step was repeated two times.
8. Additional $10-\mathrm{min}$ washes were performed in 10 ml of Wash Solution 2 ( 0.1 X SSC , $0.1 \% \mathrm{SDS}$ ) with continuous agitation at $65^{\circ} \mathrm{C}$.
9. Using forceps, the cDNA array was removed from the container, briefly washed in 0 . IXSSC and excess buffer was removed from surface by centrifugation in a Beckman CS-6R centrifuge at 2000 rpm .
10. Glass arrays were scanned using a custom-built laser scanner equipped by green (Cy3 chanel) and red ( Cy5 chanel) solid state laser built in UCLA. Images were scanned at a resolution of $20 \mu \mathrm{~m}$ per pixel.

It is evident from the above results and discussion that the subject invention provides a rapid, high throughput means to simply and quickly obtain a broad-scale screening of gene expression in a variety of different samples. Only simple hybridization protocols need be employed with the subject arrays, and signals can be detected using any convenient and readily available detection device. Despite their simplicity, assays conducted with the
subject arrays yield a large amount of information regarding the expression of numerous different and important genes in a particular sample at substantially the same time, and thus have use in many different types of applications, including drug discovery and characterization, disease research, and the like.

All publications and patent applications cited in this specification are herein incorporated by reference as if each individual publication or patent application were specifically and individually indicated to be incorporated by reference. The citation of any publication is for its disclosure prior to the filing date and should not be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention.

Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it is readily apparent to those of ordinary skill in the art in light of the teachings of this invention that certain changes and modifications may be made thereto without departing from the spirit or scope of the appended claims.

## THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. An array comprising a plurality of polynucleotide probe spots stably associated with the surface of a solid support, wherein each polynucleotide probe spot is made up of unique polynucleotides each having a length that does not exceed about 1000 nt , wherein each of said polynucleotides has a known sequence of a known gene whose coding region has been fully sequenced, where all of the unique polynucleotides on said array correspond to genes of a specific type.
2. The array according to Claim 1, wherein said polynucleotides of said array have an average length of from 120 to 1000 nt .
3. The array according to Claim 1 or 2 , wherein said polynucleotides of said array do not exceed about 700 nt in length.
4. The array according to any one of the Claims 1 to 3, wherein each of said unique polynucleotides does not cross hybridize with the polynucleotides of any other polynucleotide probe concentration on the array.
5. The array according to any one of the Claims 1 to 4 , wherein said polynucleotide probe composition comprises a population of single stranded identical polynucleotides.
6. The array according to any one of the claims 1 to 4 , wherein said polynucleotide probe composition comprises a population of two different complementary single stranded polynucleotides.
7. The array according to any of the preceding claims, wherein the density of spots on said array does not exceed about $500 / \mathrm{cm}^{2}$.
8. The array according to any of the preceding claims, wherein the number of spots on said array ranges from about 50 to 1000.
9. The array according to any of the preceding claims, wherein said array is selected from the group consisting of a human array, a mouse array, a cancer array, an apoptosis array, a human stress array, an oncogene/tumor suppressor array, a cell-cell interaction array, a cytokine and cytokine receptor array, a rat array, a blood array, a mouse stress array, and a neuroarray
10. The array according to any of the preceding claims, wherein said solid support is flexible.
11. The array according to any of the preceding claims, wherein said solid support is rigid.
12. The array according to any of the preceding claims, wherein said polynucleotide probes of said array are those listed in a table selected from the group consisting of: Table 1, Table 2, Table 3, Table 4, Table 5, Table 6, Table 7 and Table 8.
13. A method of preparing an array according to any of the preceding claims, said method comprising:
enzymatically generating said unique polynucleotides; and stably associating said enzymatically generated, complementary, unique polynucleotides on the surface of said solid support.
array according to any of claims 1 to 12 under hybridization conditions sufficient to produce a hybridization pattern; and detecting said hybridization pattern.
14. The method according to claim 19, wherein said method further comprises washing said array prior to said detection step.
15. The method according to claims 19 or 20 , wherein said method further comprises preparing said labeled target polynucleotide sample.
16. The method according to claim 21, wherein said preparation comprises:
obtaining a sample of nucleic acids from a physiological source; and
generating a population of labeled nucleic acids from the nucleic acids sample by using a set of a representative number of distinct gene specific primers according to any of claims 14 to 18 ;
whereby said labeled target polynucleotide sample is produced.
17. The method according to claims 21 or 22, wherein said preparing comprises conjugating a detectable label to a functionalized target polynucleotide.
18. The method according to any of the claims 19 to 23 , wherein said method further comprises:
generating a second hybridization pattern; and comparing said hybridization patterns.
19. The method according to claim 24, wherein said hybridization patterns are generated on the same array.
20. The method according to claim 24, wherein said hybridization patterns are generated on different arrays.
21. A kit for use in a hybridization assay, said kit comprising: an array according to any of claims 1 to 12 .
22. The kit according to claim 27, wherein said kit further comprises reagents for generating a labeled target polynucleotide sample.
23. The kit according to claim 28, wherein said reagents comprise a set of a representational number of gene specific primers according to any of claims 14 to 18.
24. A kit for use in detecting the differential expression of genes of a plurality of physiological sources, the kit comprising:
a set of a representative number of distinct gene specific primers according to any of claims 14 to 18 .
25. An array according to any one of claims 1 to 12 , substantially as hereinbefore described with particular reference to the examples.
26. A method according to any one of claims 13 or 19 to 26 , substantially as hereinbefore described with particular reference to the examples.
27. A composition of gene primers according to any one of claims 14 to 18 , substantially as hereinbefore described with particular reference to the examples.
28. A kit according to any one of claims 27 to 30, substantially as hereinbefore described with particular reference to the examples.

Dated this tenth day of August 2001

Clontech Laboratories, Inc
Patent Attorneys for the Applicant:

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[^0]:    $2 \mu \mathrm{I} \quad 5 \mathrm{X}$ First-strand buffer ( $250 \mu \mathrm{M}$ Tris- HCl pH 8.3 : $375 \mathrm{mM} \mathrm{KCl} ; 15 \mathrm{mM} \mathrm{MgCl}{ }_{2}$ )
    $|\mu| \quad 10 X d N T P$ mix ( $500 \mu \mathrm{M}$ dGTP, $500 \mu \mathrm{M} \mathrm{dCTP}, 500 \mu \mathrm{M} \mathrm{dTTP}, 5 \mu \mathrm{MdATP}$ )
    $4 \mu \mathrm{l} \quad\left[\alpha^{-3} \mathrm{P}\right.$ P]dATP (Amersham, $3000 \mathrm{Ci} / \mathrm{mmol}, 10 \mathrm{mCi} / \mathrm{ml}$ )
    $1 \mu \mid \quad$ MMLV reverse transcriptase (Amersham. 200 units $/ \mu \mathrm{l}$ )

