

METHOD TO USE GENE EXPRESSION TO DETERMINE LIKELIHOOD OF CLINICAL OUTCOME OF RENAL CANCER

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application claims priority benefit of U.S. provisional application serial no. 61/294,038, filed January 11, 2010 and priority benefit of U.S. provisional application serial no. 61/346,230, filed May 19, 2010, each of which applications are incorporated herein in their entireties.

TECHNICAL FIELD

[0002] The present disclosure relates to molecular diagnostic assays that provide information concerning prognosis in renal cancer patients.

INTRODUCTION

[0003] Each year in the United States there are approximately 51,000 cases of renal cell carcinoma (kidney cancer) and upper urinary tract cancer, resulting in more than 12,900 deaths. These tumors account for approximately 3% of adult malignancies. Renal cell carcinoma (RCC) represents about 3 percent of all cancers in the United States. Predictions for the United States for the year 2007 were that 40,000 new patients would be diagnosed with RCC and that 13,000 would die from this disease.

[0004] The clinical outcome for a renal cell carcinoma patient depends largely on the aggressiveness of their particular cancer. Surgical resection is the most common treatment for this disease as systemic therapy has demonstrated only limited effectiveness. However, approximately 30% of patients with localized tumors will experience a relapse following surgery, and only 40% of all patients with renal cell carcinoma survive for 5 years.

[0005] In the US, the number of adjuvant treatment decisions that will be made by patients with early stage renal cell carcinoma in 2005 exceeded 25,000. The rates in the European Union are expected to be similar. Physicians require prognostic information to help them make informed treatment decisions for patients with renal cell carcinoma and recruit appropriate high-risk patients for clinical trials. Surgeons must decide how much kidney and surrounding tissue to remove based, in part, on predicting the aggressiveness of a particular tumor. Today, cancer tumors are generally classified based on clinical and pathological features, such as stage, grade, and the presence of necrosis. These designations are made by applying

standardized criteria, the subjectivity of which has been demonstrated by a lack of concordance amongst pathology laboratories.

SUMMARY

[0006] The present disclosure provides biomarkers, the expression of which has prognostic value in renal cancer.

BRIEF DESCRIPTION OF THE DRAWINGS

[0007] Figures 1a-1c: Consistency between Stage I and III for exemplary genes associated with RFI

[0008] Figure 2: Consistent results across endpoints (OS and RFI) for exemplary genes

[0009] Figure 3: Kaplan-Meier curve: Recurrence Free Interval (RFI) by Cleveland Clinic Foundation (CCF) histologic necrosis

[0010] Figure 4: Performance of Mayo prognostic tool applied to CCF data

[0011] Figure 5: Example of using one gene to improve estimate: EMCN in addition to Mayo Criteria

[0012] Figure 6: Example of using one gene to improve estimate: AQP1 in addition to Mayo Criteria

[0013] Figure 7: Example of using one gene to improve estimate: PPAP2B in addition to Mayo Criteria

DETAILED DESCRIPTION

DEFINITIONS

[0014] Unless defined otherwise, technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Singleton *et al.*, Dictionary of Microbiology and Molecular Biology 2nd ed., J. Wiley & Sons (New York, NY 1994), and March, Advanced Organic Chemistry Reactions, Mechanisms and Structure 4th ed., John Wiley & Sons (New York, NY 1992), provide one skilled in the art with a general guide to many of the terms used in the present application.

[0015] One skilled in the art will recognize many methods and materials similar or equivalent to those described herein, which could be used in the practice of the present invention. Indeed, the present invention is in no way limited to the methods and materials described. For purposes of the present invention, the following terms are defined below.

[0016] The term “tumor” is used herein to refer to all neoplastic cell growth and proliferation, and all pre-cancerous and cancerous cells and tissues. The term “primary tumor” is used herein to refer to a tumor that is at the original site where it first arose. For example, a primary renal cell carcinoma tumor is one that arose in the kidney. The term “metastatic tumor” is used herein to refer to a tumor that develops away from the site of origin. For example, renal cell carcinoma metastasis most commonly affects the spine, ribs, pelvis, and proximal long bones.

[0017] The terms "cancer" and "carcinoma" refer to or describe the physiological condition in mammals that is typically characterized by unregulated cell growth. The pathology of cancer includes, for example, abnormal or uncontrollable cell growth, metastasis, interference with the normal functioning of neighboring cells, release of cytokines or other secretory products at abnormal levels, suppression or aggravation of inflammatory or immunological response, neoplasia, premalignancy, malignancy, invasion of surrounding or distant tissues or organs, such as lymph nodes, blood vessels, etc.

[0018] As used herein, the terms “renal cancer” or “renal cell carcinoma” refer to cancer that has arisen from the kidney.

[0019] The terms “renal cell cancer” or “renal cell carcinoma” (RCC), as used herein, refer to cancer which originates in the lining of the proximal convoluted tubule. More specifically, RCC encompasses several relatively common histologic subtypes: clear cell renal cell carcinoma, papillary (chromophil), chromophobe, collecting duct carcinoma, and medullary carcinoma. Further information about renal cell carcinoma may be found in Y. Thyaviahally, et al., *Int Semin Surg Oncol* 2:18 (2005), the contents of which are incorporated by reference herein. Clear cell renal cell carcinoma (ccRCC) is the most common subtype of RCC. Incidence of ccRCC is increasing, comprising 80% of localized disease and more than 90% of metastatic disease.

[0020] The staging system for renal cell carcinoma is based on the degree of tumor spread beyond the kidney. According to the tumor, node, metastasis (TNM) staging system of the American Joint Committee on Cancer (AJCC) (Greene, et al., *AJCC Cancer Staging Manual*, pp. 323-325 (6th Ed. 2002), the various stages of renal cell carcinoma are provided below. “Increased stage” as used herein refers to classification of a tumor at a stage that is more advanced, e.g., Stage 4 is an increased stage relative to Stages 1, 2, and 3.

<u>Description of RCC Stages</u>
<u>Stages for Renal Cell Carcinoma</u>
Stage 1: T1, N0, M0
Stage 2: T2, N0, M0
Stage 3: T1-T2, N1, M0; T3, N0-1, M0; T3a, N0-1, M0; T3b, N0-1, M0; and T3c, N0-1, M0
Stage 4: T4, N0-1, M0; Any T, N2, M0; and Any T, any N, M1
<u>Primary tumor (T)</u>
TX: Primary tumor cannot be assessed
T0: No evidence of primary tumor
T1: Tumor 7 cm or less in greatest dimension and limited to the kidney
T1a: Tumor 4 cm or less in greatest dimension and limited to the kidney
T1b: Tumor larger than 4 cm but 7 cm or less in greatest dimension and limited to the kidney
T2: Tumor larger than 7 cm in greatest dimension and limited to the kidney
T3: Tumor extends into major veins or invades adrenal gland or perinephric tissues but not beyond Gerota fascia
T3a: Tumor directly invades adrenal gland or perirenal and/or renal sinus fat but not beyond Gerota fascia
T3b: Tumor grossly extends into the renal vein or its segmental (i.e., muscle-containing) branches, or it extends into the vena cava below the diaphragm
T3c: Tumor grossly extends into the vena cava above the diaphragm or invades the wall of the vena cava
T4: Tumor invades beyond Gerota fascia
<u>Regional lymph nodes (N)</u>
NX: Regional lymph nodes cannot be assessed
N0: No regional lymph node metastasis
N1: Metastasis in a single regional lymph node
N2: Metastasis in more than one regional lymph node
<u>Distant metastasis (M)</u>
MX: Distant metastasis cannot be assessed
M0: No distant metastasis
M1: Distant metastasis

[0021] The term “early stage renal cancer”, as used herein, refers to Stages 1-3, as defined in the American Joint Committee on Cancer (AJCC) Cancer Staging Manual, pp. 323-325 (6th Ed. 2002).

[0022] Reference to tumor “grade” for renal cell carcinoma as used herein refers to a grading system based on microscopic appearance of tumor cells. According to the TNM staging system of the AJCC, the various grades of renal cell carcinoma are:

- GX (grade of differentiation cannot be assessed);
- G1 (well differentiated);

G2 (moderately differentiated); and

G3-G4 (poorly differentiated/undifferentiated).

“Increased grade” as used herein refers to classification of a tumor at a grade that is more advanced, e.g., Grade 4 (G4) 4 is an increased grade relative to Grades 1, 2, and 3. Tumor grading is an important prognostic factor in renal cell carcinoma. H. Rauschmeier, et al., *World J Urol* 2:103-108 (1984).

[0023] The terms “necrosis” or “histologic necrosis” as used herein refer to the death of living cells or tissues. The presence of necrosis may be a prognostic factor in cancer. For example, necrosis is commonly seen in renal cell carcinoma (RCC) and has been shown to be an adverse prognostic factor in certain RCC subtypes. V. Foria, et al., *J Clin Pathol* 58(1):39-43 (2005).

[0024] The terms “nodal invasion” or “node-positive (N+)” as used herein refer to the presence of cancer cells in one or more lymph nodes associated with the organ (e.g., drain the organ) containing a primary tumor. Nodal invasion is part of tumor staging for most cancers, including renal cell carcinoma.

[0025] The term “prognostic gene,” when used in the single or plural, refers to a gene, the expression level of which is correlated with a good or bad prognosis for a cancer patient. A gene may be both a prognostic and predictive gene, depending on the association of the gene expression level with the corresponding endpoint.

[0026] The terms “correlated” and “associated” are used interchangeably herein to refer to the strength of association between two measurements (or measured entities). The disclosure provides genes and gene subsets, the expression levels of which are associated with a particular outcome measure, such as between the expression level of a gene and the likelihood of a recurrence event or relapse. For example, the increased expression level of a gene may be positively correlated (positively associated) with an increased likelihood of good clinical outcome for the patient, such as a decreased likelihood of recurrence of cancer. Such a positive correlation may be demonstrated statistically in various ways, e.g. by a hazard ratio less than 1.0. In another example, the increased expression level of a gene may be negatively correlated (negatively associated) with an increased likelihood of good clinical outcome for the patient. In that case, for example, a patient with a high expression level of a gene may have an increased likelihood of recurrence of the cancer. Such a negative correlation could indicate that the patient

with a high expression level of a gene likely has a poor prognosis, or might respond poorly to a chemotherapy, and this may be demonstrated statistically in various ways, e.g., a hazard ratio greater than 1.0.

[0027] “Co-expression” is used herein to refer to strength of association between the expression levels of two different genes that are biologically similar, such that expression level of a first gene may be substituted with an expression level of a second gene in a given analysis in view of their correlation of expression. Such co-expressed genes (or correlated expression) indicates that these two genes are substitutable in an expression algorithm, for example, if a first gene is highly correlated, positively or negatively, with increased likelihood of a good clinical outcome for renal cell carcinoma, then the second co-expressed gene also correlates, in the same direction as the first gene, with the same outcome. Pairwise co-expression may be calculated by various methods known in the art, e.g., by calculating Pearson correlation coefficients or Spearman correlation coefficients or by clustering methods. The methods described herein may incorporate one or more genes that co-express, with a Pearson correlation co-efficient of at least 0.5. Co-expressed gene cliques may also be identified using graph theory. An analysis of co-expression may be calculated using normalized or standardized and normalized expression data.

[0028] The terms “prognosis” and “clinical outcome” are used interchangeably herein to refer to an estimate of the likelihood of cancer-attributable death or progression, including recurrence, and metastatic spread of a neoplastic disease, such as renal cell carcinoma. The terms “good prognosis” or “positive clinical outcome” mean a desired clinical outcome. For example, in the context of renal cell carcinoma, a good prognosis may be an expectation of no local recurrences or metastasis within two, three, four, five or more years of the initial diagnosis of renal cell carcinoma. The terms “poor prognosis” or “negative clinical outcome” are used herein interchangeably to mean an undesired clinical outcome. For example, in the context of renal cell carcinoma, a poor prognosis may be an expectation of a local recurrence or metastasis within two, three, four, five or more years of the initial diagnosis of renal cell carcinoma.

[0029] The term “predictive gene” is used herein to refer to a gene, the expression of which is correlated, positively or negatively, with likelihood of beneficial response to treatment.

[0030] A “clinical outcome” can be assessed using any endpoint, including, without limitation, (1) aggressiveness of tumor growth (e.g., movement to higher stage); (2) metastasis; (3) local recurrence; (4) increase in the length of survival following treatment; and/or (5)

decreased mortality at a given point of time following treatment. Clinical response may also be expressed in terms of various measures of clinical outcome. Clinical outcome can also be considered in the context of an individual's outcome relative to an outcome of a population of patients having a comparable clinical diagnosis, and can be assessed using various endpoints such as an increase in the duration of Recurrence-Free interval (RFI), an increase in the duration of Overall Survival (OS) in a population, an increase in the duration of Disease-Free Survival (DFS), an increase in the duration of Distant Recurrence-Free Interval (DRFI), and the like.

[0031] The term "treatment", as used herein, refers to therapeutic compounds administered to patients to cease or reduce proliferation of cancer cells, shrink the tumor, avoid progression and metastasis, or cause primary tumor or metastases regression. Examples of treatment include, for example, cytokine therapy, progestational agents, anti-angiogenic therapy, hormonal therapy, and chemotherapy (including small molecules and biologics).

[0032] The terms "surgery" or "surgical resection" are used herein to refer to surgical removal of some or all of a tumor, and usually some of the surrounding tissue. Examples of surgical techniques include laproscopic procedures, biopsy, or tumor ablation, such as cryotherapy, radio frequency ablation, and high intensity ultrasound. In cancer patients, the extent of tissue removed during surgery depends on the state of the tumor as observed by a surgeon. For example, a partial nephrectomy indicates that part of one kidney is removed; a simple nephrectomy entails removal of all of one kidney; a radical nephrectomy, all of one kidney and neighboring tissue (e.g., adrenal gland, lymph nodes) removed; and bilateral nephrectomy, both kidneys removed.

[0033] The terms "recurrence" and "relapse" are used herein, in the context of potential clinical outcomes of cancer, to refer to a local or distant metastases. Identification of a recurrence could be done by, for example, CT imaging, ultrasound, arteriogram, or X-ray, biopsy, urine or blood test, physical exam, or research center tumor registry.

[0034] The term "recurrence-free interval" as used herein refers to the time from surgery to first recurrence or death due to recurrence of renal cancer. Losses to follow-up, second primary cancers, other primary cancers, and deaths prior to recurrence are considered censoring events.

[0035] The term “overall survival” is defined as the time from surgery to death from any cause. Losses to follow-up are considered censoring events. Recurrences are ignored for the purposes of calculating overall survival (OS).

[0036] The term “disease-free survival” is defined as the time from surgery to first recurrence or death from any cause, whichever occurs first. Losses to follow-up are considered censoring events.

[0037] The term “Hazard Ratio (HR)” as used herein refers to the effect of an explanatory variable on the hazard or risk of an event (i.e. recurrence or death). In proportional hazards regression models, the HR is the ratio of the predicted hazard for two groups (e.g. patients with or without necrosis) or for a unit change in a continuous variable (e.g. one standard deviation change in gene expression).

[0038] The term “Odds Ratio (OR)” as used herein refers to the effect of an explanatory variable on the odds of an event (e.g. presence of necrosis). In logistic regression models, the OR is the ratio of the predicted odds for a unit change in a continuous variable (e.g. one standard deviation change in gene expression).

[0039] The term “prognostic clinical and/or pathologic covariate” as used herein refers to clinical and/or prognostic covariates that are significantly associated ($p \leq 0.05$) with clinical outcome. For example, prognostic clinical and pathologic covariates in renal cell carcinoma include tumor stage (e.g. size, nodal invasion, etc.), and grade (e.g., Fuhrman grade), histologic necrosis, and gender.

[0040] The term “proxy gene” refers to a gene, the expression of which is correlated (positively or negatively) with one or more prognostic clinical and/or pathologic covariates. The expression level(s) of one or more proxy genes may be used instead of, or in addition to, classification of a tumor by physical or mechanical examination in a pathology laboratory.

[0041] The term “microarray” refers to an ordered arrangement of hybridizable array elements, preferably polynucleotide probes, on a substrate.

[0042] The term “polynucleotide,” when used in singular or plural, generally refers to any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. Thus, for instance, polynucleotides as defined herein include, without limitation, single- and double-stranded DNA, DNA including single- and double-stranded regions, single- and double-stranded RNA, and RNA including single- and double-stranded

regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or include single- and double-stranded regions. In addition, the term “polynucleotide” as used herein refers to triple-stranded regions comprising RNA or DNA or both RNA and DNA. The strands in such regions may be from the same molecule or from different molecules. The regions may include all of one or more of the molecules, but more typically involve only a region of some of the molecules. One of the molecules of a triple-helical region often is an oligonucleotide. The term “polynucleotide” specifically includes DNAs (e.g., cDNAs) and RNAs that contain one or more modified bases. Thus, DNAs or RNAs with backbones modified for stability or for other reasons are “polynucleotides” as that term is intended herein. Moreover, DNAs or RNAs comprising unusual bases, such as inosine, or modified bases, such as tritiated bases, are included within the term “polynucleotides” as defined herein. In general, the term “polynucleotide” embraces all chemically, enzymatically and/or metabolically modified forms of unmodified polynucleotides, as well as the chemical forms of DNA and RNA characteristic of viruses and cells, including simple and complex cells.

[0043] The term “oligonucleotide” refers to a relatively short polynucleotide, including, without limitation, single-stranded deoxyribonucleotides, single- or double-stranded ribonucleotides, RNA:DNA hybrids and double-stranded DNAs. Oligonucleotides, such as single-stranded DNA probe oligonucleotides, are often synthesized by chemical methods, for example using automated oligonucleotide synthesizers that are commercially available. However, oligonucleotides can be made by a variety of other methods, including *in vitro* recombinant DNA-mediated techniques and by expression of DNAs in cells and organisms.

[0044] The term “expression level” as applied to a gene refers to the normalized level of a gene product.

[0045] The terms “gene product” or “expression product” are used herein interchangeably to refer to the RNA transcription products (RNA transcript) of a gene, including mRNA, and the polypeptide translation product of such RNA transcripts. A gene product can be, for example, an unspliced RNA, an mRNA, a splice variant mRNA, a microRNA, a fragmented RNA, a polypeptide, a post-translationally modified polypeptide, a splice variant polypeptide, etc.

[0046] “Stringency” of hybridization reactions is readily determinable by one of ordinary skill in the art, and generally is an empirical calculation dependent upon probe length, washing

temperature, and salt concentration. In general, longer probes require higher temperatures for proper annealing, while shorter probes need lower temperatures. Hybridization generally depends on the ability of denatured DNA to re-anneal when complementary strands are present in an environment below their melting temperature. The higher the degree of desired homology between the probe and hybridizable sequence, the higher the relative temperature that can be used. As a result, it follows that higher relative temperatures would tend to make the reaction conditions more stringent, while lower temperatures less so. For additional details and explanation of stringency of hybridization reactions, see Ausubel et al., *Current Protocols in Molecular Biology*, (Wiley Interscience, 1995).

[0047] "Stringent conditions" or "high stringency conditions", as defined herein, typically: (1) employ low ionic strength solutions and high temperature for washing, for example 0.015 M sodium chloride/0.0015 M sodium citrate/0.1% sodium dodecyl sulfate at 50°C; (2) employ during hybridization a denaturing agent, such as formamide, for example, 50% (v/v) formamide with 0.1% bovine serum albumin/0.1% Ficoll/0.1% polyvinylpyrrolidone/50mM sodium phosphate buffer at pH 6.5 with 750 mM sodium chloride, 75 mM sodium citrate at 42°C; or (3) employ 50% formamide, 5 x SSC (0.75 M NaCl, 0.075 M sodium citrate), 50 mM sodium phosphate (pH 6.8), 0.1% sodium pyrophosphate, 5 x Denhardt's solution, sonicated salmon sperm DNA (50 µg/ml), 0.1% SDS, and 10% dextran sulfate at 42°C, with washes at 42°C in 0.2 x SSC (sodium chloride/sodium citrate) and 50% formamide at 55°C, followed by a high-stringency wash consisting of 0.1 x SSC containing EDTA at 55°C.

[0048] "Moderately stringent conditions" may be identified as described by Sambrook et al., *Molecular Cloning: A Laboratory Manual* (Cold Spring Harbor Press, 1989), and include the use of washing solution and hybridization conditions (e.g., temperature, ionic strength and %SDS) less stringent than those described above. An example of moderately stringent condition is overnight incubation at 37°C in a solution comprising: 20% formamide, 5 x SSC (150 mM NaCl, 15 mM trisodium citrate), 50 mM sodium phosphate (pH 7.6), 5 x Denhardt's solution, 10% dextran sulfate, and 20 mg/ml denatured sheared salmon sperm DNA, followed by washing the filters in 1 x SSC at about 37-50°C. The skilled artisan will recognize how to adjust the temperature, ionic strength, etc. as necessary to accommodate factors such as probe length and the like.

[0049] In the context of the present invention, reference to "at least one," "at least two," "at least five," etc. of the genes listed in any particular gene set means any one or any and all combinations of the genes listed.

[0050] The terms "splicing" and "RNA splicing" are used interchangeably and refer to RNA processing that removes introns and joins exons to produce mature mRNA with continuous coding sequence that moves into the cytoplasm of an eukaryotic cell.

[0051] In theory, the term "exon" refers to any segment of an interrupted gene that is represented in a mature RNA product (B. Lewin, *Genes IV* (Cell Press, 1990)). In theory the term "intron" refers to any segment of DNA that is transcribed but removed from within the transcript by splicing together the exons on either side of it. Operationally, exon sequences occur in the mRNA sequence of a gene as defined by Ref. SEQ ID numbers. Operationally, intron sequences are the intervening sequences within the genomic DNA of a gene, bracketed by exon sequences and usually having GT and AG splice consensus sequences at their 5' and 3' boundaries.

[0052] A "computer-based system" refers to a system of hardware, software, and data storage medium used to analyze information. The minimum hardware of a patient computer-based system comprises a central processing unit (CPU), and hardware for data input, data output (e.g., display), and data storage. An ordinarily skilled artisan can readily appreciate that any currently available computer-based systems and/or components thereof are suitable for use in connection with the methods of the present disclosure. The data storage medium may comprise any manufacture comprising a recording of the present information as described above, or a memory access device that can access such a manufacture.

[0053] To "record" data, programming or other information on a computer readable medium refers to a process for storing information, using any such methods as known in the art. Any convenient data storage structure may be chosen, based on the means used to access the stored information. A variety of data processor programs and formats can be used for storage, e.g. word processing text file, database format, etc.

[0054] A "processor" or "computing means" references any hardware and/or software combination that will perform the functions required of it. For example, a suitable processor may be a programmable digital microprocessor such as available in the form of an electronic controller, mainframe, server or personal computer (desktop or portable). Where the processor is programmable, suitable programming can be communicated from a remote location to the

processor, or previously saved in a computer program product (such as a portable or fixed computer readable storage medium, whether magnetic, optical or solid state device based). For example, a magnetic medium or optical disk may carry the programming, and can be read by a suitable reader communicating with each processor at its corresponding station.

[0055] The present disclosure provides methods for assessing a patient's risk of recurrence of cancer, which methods comprise assaying an expression level of at least one gene, or its gene product, in a biological sample obtained from the patient. In some embodiments, the biological sample can be a tumor sample obtained from the kidney, or surrounding tissues, of the patient. In other embodiments, the biological sample is obtained from a bodily fluid, such as blood or urine.

[0056] The present disclosure provides genes useful in the methods disclosed herein. The genes are listed in Tables 3a and 3b, wherein increased expression of genes listed in Table 3a is significantly associated with a lower risk of cancer recurrence, and increased expression of genes listed in Table 3b is significantly associated with a higher risk of cancer recurrence. In some embodiments, a co-expressed gene may be used in conjunction with, or substituted for, a gene listed in Tables 3a or 3b with which it co-expresses.

[0057] The present disclosure further provides genes significantly associated, positively or negatively, with renal cancer recurrence after adjusting for clinical/pathologic covariates (stage, tumor grade, tumor size, nodal status, and presence of necrosis). For example, Table 8a lists genes wherein increased expression is significantly associated with lower risk of renal cancer recurrence after adjusting for clinical/pathologic covariates, and Table 8b lists genes wherein increased expression is significantly associated with a higher risk of renal cancer recurrence after adjusting for clinical/pathologic covariates. Of those genes listed in Tables 8a and 8b, 16 genes with significant association, positively or negatively, with good prognosis after adjusting for clinical/pathologic covariates and controlling the false discovery rate at 10% are listed in Table 9. One or more of these genes may be used separately, or in addition to, at least one of the genes listed in Tables 3a and 3b, to provide information concerning a patient's prognosis.

[0058] The present disclosure also provides proxy genes that are useful for assessing the status of clinical and/or pathologic covariates for a cancer patient. Proxy genes for tumor stage are listed in Tables 4a and 4b, wherein increased expression of genes listed in Table 4a is

significantly associated with higher tumor stage, and increased expression of genes listed in Table 4b is significantly associated with lower tumor stage. Proxy genes for tumor grade are listed in Tables 5a and 5b wherein increased expression of genes listed in Table 5a is significantly associated with higher tumor grade, and increased expression of genes listed in Table 5b is significantly associated with lower tumor grade. Proxy genes for the presence of necrosis are listed in Tables 6a and 6b, wherein expression of genes listed in Table 6a is significantly associated with the presence of necrosis, and increased expression of genes listed in Table 6b is significantly associated with the absence of necrosis. Proxy genes for nodal involvement are listed in Tables 7a and 7b wherein higher expression of genes listed in Table 7a are significantly associated with the presence of nodal invasion, and increased expression of genes listed in Table 7b are significantly associated with the absence of nodal invasion. One or more proxy genes may be used separately, or in addition to, at least one of the genes listed in Tables 3a and 3b, to provide information concerning a patient's prognosis. In some embodiments, at least two of the following proxy genes are used to provide information concerning the patient's prognosis: TSPAN7, TEK, LDB2, TIMP3, SHANK3, RGS5, KDR, SDPR, EPAS1, ID1, TGFBR2, FLT4, SDPR, ENDRB, JAG1, DLC1, and KL. In some embodiments, a co-expressed gene may be used in conjunction with, or substituted for, a proxy gene with which it co-expresses.

[0059] The present disclosure also provides sets of genes in biological pathways that are useful for assessing the likelihood that a cancer patient is likely to have a positive clinical outcome, which sets of genes are referred to herein as "gene subsets". The gene subsets include angiogenesis, immune response, transport, cell adhesion/extracellular matrix, cell cycle, and apoptosis. In some embodiments, the angiogenesis gene subset includes ADD1, ANGPTL3, APOLD1, CEACAM1, EDNRB, EMCN, ENG, EPAS1, FLT1, JAG1, KDR, KL, LDB2, NOS3, NUDT6, PPAP2B, PRKCH, PTPRB, RGS5, SHANK3, SNRK, TEK, ICAM2, and VCAM1; the immune response gene subset includes CCL5, CCR7, CD8A, CX3CL1, CXCL10, CXCL9, HLA-DPB1, IL6, IL8, and SPP1, and; the transport gene subset includes AQP1 and SGK1; the cell adhesion/extracellular matrix gene subset includes ITGB1, A2M, ITGB5, LAMB1, LOX, MMP14, TGFBR2, TIMP3, and TSPAN7; the cell cycle gene subset includes BUB1, C13orf15, CCNB1, PTTG1, TPX2, LMNB1, and TUBB2A; the apoptosis gene subset includes CASP10; the early response gene subset includes EGR1 and CYR61; the metabolic signaling gene subset

includes CA12, ENO2, UGCG, and SDPR; and the signaling gene subset includes ID1 and MAP2K3.

[0060] The present disclosure also provides genes in biological pathways targeted by chemotherapy that are correlated, positively or negatively, to a risk of cancer recurrence. These genes include KIT, PDGFA, PDGFB, PDGFC, PDGFD, PDGFRb, KRAS, RAF1, MTOR, HIF1AN, VEGFA, VEGFB, and FLT4. In some embodiments, the chemotherapy is cytokine and/or anti-angiogenic therapy. In other embodiments, the chemotherapy is sunitinib, sorafenib, temsirolimus, bevacizumab, everolimus, and/or pazopanib.

[0061] In some embodiments, a co-expressed gene may be used in conjunction with, or substituted for, a gene with which it co-expresses.

[0062] In some embodiments, the cancer is renal cell carcinoma. In other embodiments, the cancer is clear cell renal cell carcinoma (ccRCC), papillary, chromophobe, collecting duct carcinoma, and/or medullary carcinoma.

[0063] Various technological approaches for determination of expression levels of the disclosed genes are set forth in this specification, including, without limitation, reverse-transcription polymerase chain reaction (RT-PCR), microarrays, high-throughput sequencing, serial analysis of gene expression (SAGE), and Digital Gene Expression (DGE), which will be discussed in detail below. In particular aspects, the expression level of each gene may be determined in relation to various features of the expression products of the gene, including exons, introns, protein epitopes, and protein activity.

[0064] The expression levels of genes identified herein may be measured in tumor tissue. For example, the tumor tissue may be obtained upon surgical resection of the tumor, or by tumor biopsy. The expression level of the identified genes may also be measured in tumor cells recovered from sites distant from the tumor, including circulating tumor cells or body fluid (e.g., urine, blood, blood fraction, etc.).

[0065] The expression product that is assayed can be, for example, RNA or a polypeptide. The expression product may be fragmented. For example, the assay may use primers that are complementary to target sequences of an expression product and could thus measure full transcripts as well as those fragmented expression products containing the target sequence. Further information is provided in Tables A and B, which provide examples of

sequences of forward primers, reverse primers, probes and amplicons generated by use of the primers.

[0066] The RNA expression product may be assayed directly or by detection of a cDNA product resulting from a PCR-based amplification method, e.g., quantitative reverse transcription polymerase chain reaction (qRT-PCR). (See e.g., U.S. Pub. No. US2006-0008809A1.) Polypeptide expression product may be assayed using immunohistochemistry (IHC). Further, both RNA and polypeptide expression products may also be is assayed using microarrays.

CLINICAL UTILITY

[0067] Currently, of the expected clinical outcome for RCC patients is based on subjective determinations of a tumor's clinical and pathologic features. For example, physicians make decisions about the appropriate surgical procedures and adjuvant therapy based on a renal tumor's stage, grade, and the presence of necrosis. Although there are standardized measures to guide pathologists in making these decisions, the level of concordance between pathology laboratories is low. (See Al-Ayanti M et al. (2003) Arch Pathol Lab Med 127, 593-596) It would be useful to have a reproducible molecular assay for determining and/or confirming these tumor characteristics.

[0068] In addition, standard clinical criteria, by themselves, have limited ability to accurately estimate a patient's prognosis. It would be useful to have a reproducible molecular assay to assess a patient's prognosis based on the biology of his or her tumor. Such information could be used for the purposes of patient counseling, selecting patients for clinical trials (e.g., adjuvant trials), and understanding the biology of renal cell carcinoma. In addition, such a test would assist physicians in making surgical and treatment recommendations based on the biology of each patient's tumor. For example, a genomic test could stratify RCC patients based on risk of recurrence and/or likelihood of long-term survival without recurrence (relapse, metastasis, etc.). There are several ongoing and planned clinical trials for RCC therapies, including adjuvant radiation and chemotherapies. It would be useful to have a genomic test able to identify high-risk patients more accurately than standard clinical criteria, thereby further enriching an adjuvant RCC population for study. This would reduce the number of patients needed for an adjuvant trial and the time needed for definitive testing of these new agents in the adjuvant setting.

[0069] Finally, it would be useful to have a molecular assay that could predict a patient's likelihood to respond to treatment, such as chemotherapy. Again, this would facilitate individual

treatment decisions and recruiting patients for clinical trials, and increase physician and patient confidence in making healthcare decisions after being diagnosed with cancer.

REPORTING RESULTS

[0070] The methods of the present disclosure are suited for the preparation of reports summarizing the expected clinical outcome resulting from the methods of the present disclosure. A “report,” as described herein, is an electronic or tangible document that includes report elements that provide information of interest relating to a likelihood assessment and its results. A subject report includes at least a likelihood assessment, e.g., an indication as to the risk of recurrence for a subject with renal cell carcinoma. A subject report can be completely or partially electronically generated, e.g., presented on an electronic display (e.g., computer monitor). A report can further include one or more of: 1) information regarding the testing facility; 2) service provider information; 3) patient data; 4) sample data; 5) an interpretive report, which can include various information including: a) indication; b) test data, where test data can include a normalized level of one or more genes of interest, and 6) other features.

[0071] The present disclosure thus provides for methods of creating reports and the reports resulting therefrom. The report may include a summary of the expression levels of the RNA transcripts, or the expression products of such RNA transcripts, for certain genes in the cells obtained from the patient’s tumor. The report can include information relating to prognostic covariates of the patient. The report may include an estimate that the patient has an increased risk of recurrence. That estimate may be in the form of a score or patient stratifier scheme (e.g., low, intermediate, or high risk of recurrence). The report may include information relevant to assist with decisions about the appropriate surgery (e.g., partial or total nephrectomy) or treatment for the patient.

[0072] Thus, in some embodiments, the methods of the present disclosure further include generating a report that includes information regarding the patient’s likely clinical outcome, e.g. risk of recurrence. For example, the methods disclosed herein can further include a step of generating or outputting a report providing the results of a subject risk assessment, which report can be provided in the form of an electronic medium (e.g., an electronic display on a computer monitor), or in the form of a tangible medium (e.g., a report printed on paper or other tangible medium).

[0073] A report that includes information regarding the patient's likely prognosis (e.g., the likelihood that a patient having renal cell carcinoma will have a good prognosis or positive clinical outcome in response to surgery and/or treatment) is provided to a user. An assessment as to the likelihood is referred to below as a "risk report" or, simply, "risk score." A person or entity that prepares a report ("report generator") may also perform the likelihood assessment. The report generator may also perform one or more of sample gathering, sample processing, and data generation, e.g., the report generator may also perform one or more of: a) sample gathering; b) sample processing; c) measuring a level of a risk gene; d) measuring a level of a reference gene; and e) determining a normalized level of a risk gene. Alternatively, an entity other than the report generator can perform one or more sample gathering, sample processing, and data generation.

[0074] For clarity, it should be noted that the term "user," which is used interchangeably with "client," is meant to refer to a person or entity to whom a report is transmitted, and may be the same person or entity who does one or more of the following: a) collects a sample; b) processes a sample; c) provides a sample or a processed sample; and d) generates data (e.g., level of a risk gene; level of a reference gene product(s); normalized level of a risk gene for use in the likelihood assessment. In some cases, the person(s) or entity(ies) who provides sample collection and/or sample processing and/or data generation, and the person who receives the results and/or report may be different persons, but are both referred to as "users" or "clients" herein to avoid confusion. In certain embodiments, e.g., where the methods are completely executed on a single computer, the user or client provides for data input and review of data output. A "user" can be a health professional (e.g., a clinician, a laboratory technician, a physician (e.g., an oncologist, surgeon, pathologist), etc.).

[0075] In embodiments where the user only executes a portion of the method, the individual who, after computerized data processing according to the methods of the present disclosure, reviews data output (e.g., results prior to release to provide a complete report, a complete, or reviews an "incomplete" report and provides for manual intervention and completion of an interpretive report) is referred to herein as a "reviewer." The reviewer may be located at a location remote to the user (e.g., at a service provided separate from a healthcare facility where a user may be located).

[0076] Where government regulations or other restrictions apply (e.g., requirements by health, malpractice, or liability insurance), all results, whether generated wholly or partially electronically, are subjected to a quality control routine prior to release to the user.

METHODS OF ASSAYING EXPRESSION LEVELS OF A GENE PRODUCT

[0077] Numerous assay methods for measuring an expression level of a gene product are known in the art, including assay methods for measuring an expression level of a nucleic acid gene product (e.g., an mRNA), and assay methods for measuring an expression level of a polypeptide gene product.

Measuring a Level of a Nucleic Acid Gene Product

[0078] In general, methods of measuring a level of a nucleic acid gene product (e.g., an mRNA) include methods involving hybridization analysis of polynucleotides, and methods involving amplification of polynucleotides. Commonly used methods known in the art for the quantification of mRNA expression in a sample include northern blotting and *in situ* hybridization (See for example, Parker & Barnes, *Methods in Molecular Biology* 106:247-283 (1999)); RNase protection assays (Hod, *Biotechniques* 13:852-854 (1992)); and reverse transcription polymerase chain reaction (RT-PCR) (Weis *et al.*, *Trends in Genetics* 8:263-264 (1992)). Alternatively, antibodies may be employed that can recognize specific duplexes, including DNA duplexes, RNA duplexes, and DNA-RNA hybrid duplexes or DNA-protein duplexes. Representative methods for sequencing-based gene expression analysis include Serial Analysis of Gene Expression (SAGE), and gene expression analysis by massively parallel signature sequencing (MPSS).

Expression Methods Based on Hybridization

[0079] The level of a target nucleic acid can be measured using a probe that hybridizes to the target nucleic acid. The target nucleic acid could be, for example, a RNA expression product of a response indicator gene associated with response to a VEGF/VEGFR Inhibitor, or a RNA expression product of a reference gene. In some embodiments, the target nucleic acid is first amplified, for example using a polymerase chain reaction (PCR) method.

[0080] A number of methods are available for analyzing nucleic acid mixtures for the presence and/or level of a specific nucleic acid. mRNA may be assayed directly or reverse transcribed into cDNA for analysis.

[0081] In some embodiments, the method involves contacting a sample (e.g., a sample derived from a cancer cell) under stringent hybridization conditions with a nucleic acid probe and detecting binding, if any, of the probe to a nucleic acid in the sample. A variety of nucleic acid hybridization methods are well known to those skilled in the art, and any known method can be used. In some embodiments, the nucleic acid probe will be detectably labeled.

Expression Methods Based on Target Amplification

[0082] Methods of amplifying (e.g., by PCR) nucleic acid, methods of performing primers extension, and methods of assessing nucleic acids are generally well known in the art. (See e.g., Ausubel, et al, Short Protocols in Molecular Biology, 3rd ed., Wiley & Sons, 1995 and Sambrook, et al, Molecular Cloning: A Laboratory Manual, Third Edition, (2001) Cold Spring Harbor, N.Y.)

[0083] A target mRNA can be amplified by reverse transcribing the mRNA into cDNA, and then performing PCR (reverse transcription-PCR or RT-PCR). Alternatively, a single enzyme may be used for both steps as described in U.S. Pat. No. 5,322,770.

[0084] The fluorogenic 5' nuclease assay, known as the TaqMan® assay (Roche Molecular Systems, Inc.), is a powerful and versatile PCR-based detection system for nucleic acid targets. For a detailed description of the TaqMan assay, reagents and conditions for use therein, see, e.g., Holland et al., Proc. Natl. Acad. Sci., U.S.A. (1991) 88:7276-7280; U.S. Pat. Nos. 5,538,848, 5,723,591, and 5,876,930, all incorporated herein by reference in their entireties. Hence, primers and probes derived from regions of a target nucleic acid as described herein can be used in TaqMan analyses to detect a level of target mRNA in a biological sample. Analysis is performed in conjunction with thermal cycling by monitoring the generation of fluorescence signals. (TaqMan is a registered trademark of Roche Molecular Systems.)

[0085] The fluorogenic 5' nuclease assay is conveniently performed using, for example, AmpliTaq Gold® DNA polymerase, which has endogenous 5' nuclease activity, to digest an internal oligonucleotide probe labeled with both a fluorescent reporter dye and a quencher (see, Holland et al., Proc Nat Acad Sci USA (1991) 88:7276-7280; and Lee et al., Nucl. Acids Res. (1993) 21:3761-3766). Assay results are detected by measuring changes in fluorescence that occur during the amplification cycle as the fluorescent probe is digested, uncoupling the dye and quencher labels and causing an increase in the fluorescent signal that is proportional to the

amplification of target nucleic acid. (AmpliTaq Gold is a registered trademark of Roche Molecular Systems.)

[0086] The amplification products can be detected in solution or using solid supports. In this method, the TaqMan probe is designed to hybridize to a target sequence within the desired PCR product. The 5' end of the TaqMan probe contains a fluorescent reporter dye. The 3' end of the probe is blocked to prevent probe extension and contains a dye that will quench the fluorescence of the 5' fluorophore. During subsequent amplification, the 5' fluorescent label is cleaved off if a polymerase with 5' exonuclease activity is present in the reaction. Excision of the 5' fluorophore results in an increase in fluorescence that can be detected.

[0087] The first step for gene expression analysis is the isolation of mRNA from a target sample. The starting material is typically total RNA isolated from human tumors or tumor cell lines, and corresponding normal tissues or cell lines, respectively. Thus RNA can be isolated from a variety of primary tumors, including breast, lung, colon, prostate, brain, liver, kidney, pancreas, spleen, thymus, testis, ovary, uterus, head and neck, etc., tumor, or tumor cell lines, with pooled DNA from healthy donors. If the source of mRNA is a primary tumor, mRNA can be extracted, for example, from frozen or archived paraffin-embedded and fixed (e.g., formalin-fixed) tissue samples.

[0088] General methods for mRNA extraction are well known in the art and are disclosed in standard textbooks of molecular biology, including Ausubel *et al.*, *Current Protocols of Molecular Biology* (Wiley and Sons, 1997). Methods for RNA extraction from paraffin embedded tissues are disclosed, for example, M. Cronin, *Am J. Pathol* 164(1):35-42 (2004), the contents of which are incorporated herein. In particular, RNA isolation can be performed using kits and reagents from commercial manufacturers according to the manufacturer's instructions. For example, total RNA from cells in culture can be isolated using RNeasy® mini-columns (Qiagen GmbH Corp.). Other commercially available RNA isolation kits include MasterPure™ Complete DNA and RNA Purification Kit (EPICENTRE® Biotechnologies, Madison, WI), mirVana (Applied Biosystems, Inc.), and Paraffin Block RNA Isolation Kit (Ambion, Inc.). Total RNA from tissue samples can be isolated using RNA STAT-60™ (IsoTex Diagnostics, Inc., Friendswood TX). RNA prepared from tumor can be isolated, for example, by cesium chloride density gradient centrifugation. (RNeasy is a registered trademark of Qiagen GmbH

Corp.; MasterPure is a trademark of EPICENTRE Biotechnologies; RNA STAT-60 is a trademark of Tel-Test Inc.)

[0089] As RNA cannot serve as a template for PCR, the first step in gene expression profiling by RT-PCR is the reverse transcription of the RNA template into cDNA, followed by its exponential amplification in a PCR reaction. The two most commonly used reverse transcriptase enzymes are avilo myeloblastosis virus reverse transcriptase (AMV-RT) and Moloney murine leukemia virus reverse transcriptase (MMLV-RT). The reverse transcription step is typically primed using specific primers, random hexamers, or oligo-dT primers, depending on the circumstances and the goal of expression profiling. For example, extracted RNA can be reverse-transcribed using a GeneAmp® RNA PCR kit (Applied Biosystems Inc., Foster City, CA) according to the manufacturer's instructions. The derived cDNA can then be used as a template in a subsequent PCR reaction. (GeneAmp is a registered trademark of Applied Biosystems Inc.)

[0090] Although the PCR step can use a variety of thermostable DNA-dependent DNA polymerases, it typically employs the Taq DNA polymerase, which has a 5'-3' nuclease activity but lacks a 3'-5' proofreading endonuclease activity. Thus, TaqMan PCR typically utilizes the 5'-nuclease activity of Taq or Tth polymerase to hydrolyze a hybridization probe bound to its target amplicon, but any enzyme with equivalent 5' nuclease activity can be used. Two oligonucleotide primers are used to generate an amplicon. A third oligonucleotide, or probe, is designed to detect nucleotide sequence located between the two PCR primers. The probe is non-extendible by Taq DNA polymerase enzyme, and is labeled with a reporter fluorescent dye and a quencher fluorescent dye. Any laser-induced emission from the reporter dye is quenched by the quenching dye when the two dyes are located close together as they are on the probe. During the amplification reaction, the Taq DNA polymerase enzyme cleaves the probe in a template-dependent manner. The resultant probe fragments disassociate in solution, and signal from the released reporter dye is free from the quenching effect of the second fluorophore. One molecule of reporter dye is liberated for each new molecule synthesized, and detection of the unquenched reporter dye provides the basis for quantitative interpretation of the data. (TaqMan is a registered mark of Applied Biosystems.)

[0091] TaqMan RT-PCR can be performed using commercially available equipment, such as, for example, the ABI PRISM 7700® Sequence Detection System (Applied Biosystems,

Foster City, CA, USA), or the Lightcycler® (Roche Molecular Biochemicals, Mannheim, Germany). In a preferred embodiment, the 5' nuclease procedure is run on a real-time quantitative PCR device such as the ABI PRISM 7900™ Sequence Detection System™. The system consists of a thermocycler, laser, charge-coupled device (CCD), camera and computer. The system amplifies samples in a 96-well format on a thermocycler. During amplification, laser-induced fluorescent signal is collected in real-time through fiber optics cables for all 96 wells, and detected at the CCD. The system includes software for running the instrument and for analyzing the data. (PRISM 7700 is a registered trademark of Applied Biosystems; Lightcycler is a registered trademark of Roche Diagnostics GmbH LLC.)

[0092] 5'-Nuclease assay data are initially expressed as C_t , or the threshold cycle. As discussed above, fluorescence values are recorded during every cycle and represent the amount of product amplified to that point in the amplification reaction. The point when the fluorescent signal is first recorded as statistically significant is the threshold cycle (C_t).

[0093] To minimize the effect of sample-to-sample variation, quantitative RT-PCR is usually performed using an internal standard, or one or more reference genes. The ideal internal standard is expressed at a constant level among different tissues, and is unaffected by the experimental treatment. RNAs that can be used to normalize patterns of gene expression include, e.g., mRNAs for the reference genes glyceraldehyde-3-phosphate-dehydrogenase (GAPDH) and β -actin.

[0094] A more recent variation of the RT-PCR technique is the real time quantitative PCR, which measures PCR product accumulation through a dual-labeled fluorogenic probe (i.e., TaqMan® probe). Real time PCR is compatible both with quantitative competitive PCR, where internal competitor for each target sequence is used for normalization, and with quantitative comparative PCR using a normalization gene contained within the sample, or a reference gene for RT-PCR. For further details see, e.g., Held et al., Genome Research 6:986-994 (1996).

[0095] Factors considered in PCR primer design include primer length, melting temperature (T_m), and G/C content, specificity, complementary primer sequences, and 3'-end sequence. In general, optimal PCR primers are generally 17-30 bases in length, and contain about 20-80%, such as, for example, about 50-60% G+C bases. T_m 's between 50 and 80 °C, e.g., about 50 to 70 °C can be used.

[0096] For further guidelines for PCR primer and probe design see, e.g., Dieffenbach, C.W. et al., "General Concepts for PCR Primer Design" in: PCR Primer, A Laboratory Manual, Cold Spring Harbor Laboratory Press, New York, 1995, pp. 133-155; Innis and Gelfand, "Optimization of PCRs" in: *PCR Protocols, A Guide to Methods and Applications*, CRC Press, London, 1994, pp. 5-11; and Plasterer, T.N. PrimerSelect: Primer and probe design. *Methods Mol. Biol.* 70:520-527 (1997), the entire disclosures of which are hereby expressly incorporated by reference.

[0097] Other suitable methods for assaying a level of a nucleic acid gene product include, e.g., microarrays; serial analysis of gene expression (SAGE); MassARRAY® analysis; digital gene expression (DGE) (J. Marioni, *Genome Research* 18(9):1509-1517 (2008), gene expression by massively parallel signature sequencing (see, e.g., Ding and Cantor, *Proc. Nat'l Acad Sci* 100:3059-3064 (2003)); differential display (Liang and Pardee, *Science* 257:967-971 (1992)); amplified fragment length polymorphism (iAFLP) (Kawamoto et al., *Genome Res.* 12:1305-1312 (1999)); BeadArray™ technology (Illumina, San Diego, CA; Oliphant et al., *Discovery of Markers for Disease* (Supplement to *Biotechniques*), June 2002; Ferguson et al., *Analytical Chemistry* 72:5618 (2000)); BeadsArray for Detection of Gene Expression (BADGE), using the commercially available Luminex100 LabMAP system and multiple color-coded microspheres (Luminex Corp., Austin, TX) in a rapid assay for gene expression (Yang et al., *Genome Res.* 11:1888-1898 (2001)); and high coverage expression profiling (HiCEP) analysis (Fukumura et al., *Nucl. Acids. Res.* 31(16) e94 (2003)).

Introns

[0098] Assays to measure the amount of an RNA gene expression product can be targeted to intron sequences or exon sequences of the primary transcript. The amount of a spliced intron that is measured in human tissue samples is generally indicative of the amount of a corresponding exon (i.e. an exon from the same gene) present in the samples. Polynucleotides that consist of or are complementary to intron sequences can be used, e.g., in hybridization methods or amplification methods to assay the expression level of response indicator genes.

Measuring levels of a polypeptide gene product

[0099] Methods of measuring a level of a polypeptide gene product are known in the art and include antibody-based methods such as enzyme-linked immunoabsorbent assay (ELISA), radioimmunoassay (RIA), protein blot analysis, immunohistochemical analysis and the like. The

measure of a polypeptide gene product may also be measured in vivo in the subject using an antibody that specifically binds a target polypeptide, coupled to a paramagnetic label or other label used for in vivo imaging, and visualizing the distribution of the labeled antibody within the subject using an appropriate in vivo imaging method, such as magnetic resonance imaging. Such methods also include proteomics methods such as mass spectrometric methods, which are known in the art.

Methods of Isolating RNA from Body Fluids

[00100] Methods of isolating RNA for expression analysis from blood, plasma and serum (See for example, Tsui NB et al. (2002) 48,1647-53 and references cited therein) and from urine (See for example, Boom R et al. (1990) J Clin Microbiol. 28, 495-503 and reference cited therein) have been described.

Methods of Isolating RNA from Paraffin-embedded Tissue

[00101] The steps of a representative protocol for profiling gene expression using fixed, paraffin-embedded tissues as the RNA source, including mRNA isolation, purification primer extension and amplification are provided in various published journal articles. (See, e.g., T.E. Godfrey et al., J. Molec. Diagnostics 2: 84-91 (2000); K. Specht et al., Am. J. Pathol. 158: 419-29 (2001), M. Cronin, et al., Am J Pathol 164:35-42 (2004)).

MANUAL AND COMPUTER-ASSISTED METHODS AND PRODUCTS

[00102] The methods and systems described herein can be implemented in numerous ways. In one embodiment of particular interest, the methods involve use of a communications infrastructure, for example the Internet. Several embodiments are discussed below. It is also to be understood that the present disclosure may be implemented in various forms of hardware, software, firmware, processors, or a combination thereof. The methods and systems described herein can be implemented as a combination of hardware and software. The software can be implemented as an application program tangibly embodied on a program storage device, or different portions of the software implemented in the user's computing environment (e.g., as an applet) and on the reviewer's computing environment, where the reviewer may be located at a remote site associated (e.g., at a service provider's facility).

[00103] For example, during or after data input by the user, portions of the data processing can be performed in the user-side computing environment. For example, the user-side computing environment can be programmed to provide for defined test codes to denote a likelihood "risk

score,” where the score is transmitted as processed or partially processed responses to the reviewer’s computing environment in the form of test code for subsequent execution of one or more algorithms to provide a results and/or generate a report in the reviewer’s computing environment. The risk score can be a numerical score (representative of a numerical value) or a non-numerical score representative of a numerical value or range of numerical values (e.g., low, intermediate, or high).

[00104] The application program for executing the algorithms described herein may be uploaded to, and executed by, a machine comprising any suitable architecture. In general, the machine involves a computer platform having hardware such as one or more central processing units (CPU), a random access memory (RAM), and input/output (I/O) interface(s). The computer platform also includes an operating system and microinstruction code. The various processes and functions described herein may either be part of the microinstruction code or part of the application program (or a combination thereof) that is executed via the operating system. In addition, various other peripheral devices may be connected to the computer platform such as an additional data storage device and a printing device.

[00105] As a computer system, the system generally includes a processor unit. The processor unit operates to receive information, which can include test data (e.g., level of a risk gene, level of a reference gene product(s); normalized level of a gene; and may also include other data such as patient data. This information received can be stored at least temporarily in a database, and data analyzed to generate a report as described above.

[00106] Part or all of the input and output data can also be sent electronically; certain output data (e.g., reports) can be sent electronically or telephonically (e.g., by facsimile, e.g., using devices such as fax back). Exemplary output receiving devices can include a display element, a printer, a facsimile device and the like. Electronic forms of transmission and/or display can include email, interactive television, and the like. In an embodiment of particular interest, all or a portion of the input data and/or all or a portion of the output data (e.g., usually at least the final report) are maintained on a web server for access, preferably confidential access, with typical browsers. The data may be accessed or sent to health professionals as desired. The input and output data, including all or a portion of the final report, can be used to populate a patient’s medical record which may exist in a confidential database at the healthcare facility.

[00107] A system for use in the methods described herein generally includes at least one computer processor (e.g., where the method is carried out in its entirety at a single site) or at least two networked computer processors (e.g., where data is to be input by a user (also referred to herein as a "client") and transmitted to a remote site to a second computer processor for analysis, where the first and second computer processors are connected by a network, e.g., via an intranet or internet). The system can also include a user component(s) for input; and a reviewer component(s) for review of data, generated reports, and manual intervention. Additional components of the system can include a server component(s); and a database(s) for storing data (e.g., as in a database of report elements, e.g., interpretive report elements, or a relational database (RDB) which can include data input by the user and data output. The computer processors can be processors that are typically found in personal desktop computers (e.g., IBM, Dell, Macintosh), portable computers, mainframes, minicomputers, or other computing devices.

[00108] The networked client/server architecture can be selected as desired, and can be, for example, a classic two or three tier client server model. A relational database management system (RDMS), either as part of an application server component or as a separate component (RDB machine) provides the interface to the database.

[00109] In one example, the architecture is provided as a database-centric client/server architecture, in which the client application generally requests services from the application server which makes requests to the database (or the database server) to populate the report with the various report elements as required, particularly the interpretive report elements, especially the interpretation text and alerts. The server(s) (e.g., either as part of the application server machine or a separate RDB/relational database machine) responds to the client's requests.

[00110] The input client components can be complete, stand-alone personal computers offering a full range of power and features to run applications. The client component usually operates under any desired operating system and includes a communication element (e.g., a modem or other hardware for connecting to a network), one or more input devices (e.g., a keyboard, mouse, keypad, or other device used to transfer information or commands), a storage element (e.g., a hard drive or other computer-readable, computer-writable storage medium), and a display element (e.g., a monitor, television, LCD, LED, or other display device that conveys information to the user). The user enters input commands into the computer processor through an

input device. Generally, the user interface is a graphical user interface (GUI) written for web browser applications.

[00111] The server component(s) can be a personal computer, a minicomputer, or a mainframe and offers data management, information sharing between clients, network administration and security. The application and any databases used can be on the same or different servers.

[00112] Other computing arrangements for the client and server(s), including processing on a single machine such as a mainframe, a collection of machines, or other suitable configuration are contemplated. In general, the client and server machines work together to accomplish the processing of the present disclosure.

[00113] Where used, the database(s) is usually connected to the database server component and can be any device that will hold data. For example, the database can be a any magnetic or optical storing device for a computer (e.g., CDROM, internal hard drive, tape drive). The database can be located remote to the server component (with access via a network, modem, etc.) or locally to the server component.

[00114] Where used in the system and methods, the database can be a relational database that is organized and accessed according to relationships between data items. The relational database is generally composed of a plurality of tables (entities). The rows of a table represent records (collections of information about separate items) and the columns represent fields (particular attributes of a record). In its simplest conception, the relational database is a collection of data entries that "relate" to each other through at least one common field.

[00115] Additional workstations equipped with computers and printers may be used at point of service to enter data and, in some embodiments, generate appropriate reports, if desired. The computer(s) can have a shortcut (e.g., on the desktop) to launch the application to facilitate initiation of data entry, transmission, analysis, report receipt, etc. as desired.

Computer-readable storage media

[00116] The present disclosure also contemplates a computer-readable storage medium (e.g. CD-ROM, memory key, flash memory card, diskette, etc.) having stored thereon a program which, when executed in a computing environment, provides for implementation of algorithms to carry out all or a portion of the results of a response likelihood assessment as described herein. Where the computer-readable medium contains a complete program for carrying out the methods

described herein, the program includes program instructions for collecting, analyzing and generating output, and generally includes computer readable code devices for interacting with a user as described herein, processing that data in conjunction with analytical information, and generating unique printed or electronic media for that user.

[00117] Where the storage medium provides a program that provides for implementation of a portion of the methods described herein (e.g., the user-side aspect of the methods (e.g., data input, report receipt capabilities, etc.)), the program provides for transmission of data input by the user (e.g., via the internet, via an intranet, etc.) to a computing environment at a remote site. Processing or completion of processing of the data is carried out at the remote site to generate a report. After review of the report, and completion of any needed manual intervention, to provide a complete report, the complete report is then transmitted back to the user as an electronic document or printed document (e.g., fax or mailed paper report). The storage medium containing a program according to the present disclosure can be packaged with instructions (e.g., for program installation, use, etc.) recorded on a suitable substrate or a web address where such instructions may be obtained. The computer-readable storage medium can also be provided in combination with one or more reagents for carrying out response likelihood assessment (e.g., primers, probes, arrays, or other such kit components).

METHODS OF DATA ANALYSIS

Reference Normalization

[00118] In order to minimize expression measurement variations due to non-biological variations in samples, e.g., the amount and quality of expression product to be measured, raw expression level data measured for a gene product (e.g., cycle threshold (C_t) measurements obtained by qRT-PCR) may be normalized relative to the mean expression level data obtained for one or more reference genes. In one approach to normalization, a small number of genes are used as reference genes; the genes chosen for reference genes typically show a minimal amount of variation in expression from sample to sample and the expression level of other genes is compared to the relatively stable expression of the reference genes. In the global normalization approach, the expression level of each gene in a sample is compared to an average expression level in the sample of all genes in order to compare the expression of a particular gene to the total amount of material.

[00119] Unprocessed data from qRT-PCR is expressed as cycle threshold (C_t), the number of amplification cycles required for the detectable signal to exceed a defined threshold. High C_t is indicative of low expression since more cycles are required to detect the amplification product. Normalization may be carried out such that a one-unit increase in normalized expression level of a gene product generally reflects a 2-fold increase in quantity of expression product present in the sample. For further information on normalization techniques applicable to qRT-PCR data from tumor tissue, see, e.g., Silva S et al. (2006) *BMC Cancer* 6, 200; de Kok J et al. (2005) *Laboratory Investigation* 85, 154-159. Gene expression may then be standardized by dividing the normalized gene expression by the standard deviation of expression across all patients for that particular gene. By standardizing normalized gene expression the hazard ratios across genes are comparable and reflect the relative risk for each standard deviation of gene expression.

STATISTICAL ANALYSIS

[00120] One skilled in the art will recognize that variety of statistical methods are available that are suitable for comparing the expression level of a gene (or other variable) in two groups and determining the statistical significance of expression level differences that are found. (See e.g., H. Motulsky, *Intuitive Biostatistics* (Oxford University Press, 1995); D. Freedman, *Statistics* (W.W. Norton & Co, 4th Ed., 2007). For example, a Cox proportional hazards regression model may be fit to a particular clinical time-to-event endpoint (e.g., RFI, OS). One assumption of the Cox proportional hazards regression model is the proportional hazards assumption, i.e. the assumption that model effects multiply the underlying hazard. Assessments of model adequacy may be performed including, but not limited to, examination of the cumulative sum of martingale residuals. One skilled in the art would recognize that there are numerous statistical methods that may be used (e.g., Royston and Parmer (2002), smoothing spline, etc.) to fit a flexible parametric model using the hazard scale and the Weibull distribution with natural spline smoothing of the log cumulative hazards function, with effects allowed to be time-dependent. (See, P. Royston, M. Parmer, *Statistics in Medicine* 21(15):2175-2197 (2002).) The relationship between recurrence risk and (1) recurrence risk groups; and (2) clinical/pathologic covariates (e.g., tumor stage, tumor grade, presence of necrosis, lymphatic or vascular invasion, etc.) may also be tested for significance. Additional examples of models include logistic or ordinal logistic regression models in which the association between gene expression and dichotomous (for logistic) or ordinal (for ordinal logistic) clinical endpoints (i.e.

stage, necrosis, grade) may be evaluated. (See e.g., D. Hosmer and S. Lemeshow, *Applied Logistic Regression* (John Wiley and Sons, 1989).

[00121] In an exemplary embodiment, results were adjusted for multiple hypothesis tests, and allowed for a 10% false discovery rate (FDR), using Storey's procedure, and using TDRAS with separate classes (M. Crager, Gene identification using true discovery rate degree of association sets and estimates corrected for regression to the mean, *Statistics in Medicine* (published online Dec. 2009). In another embodiment, genes with significant association with RFI were identified through cross-validation techniques in which forward stepwise Cox PH regression was employed using a subset of factors identified through Principal Component Analysis (PCA).

[00122] Methods for calculating correlation coefficients, particularly the Pearson product-moment correlation coefficient are known in the art. (See e.g., J. Rodgers and W. Nicewander, *The American Statistician*, 42, 59–66 (1988); H. Motulsky, H., *Intuitive Biostatistics* (Oxford University Press, 1995). To perform particular biological processes, genes often work together in a concerted way, i.e. they are co-expressed. Co-expressed gene groups identified for a disease process like cancer can serve as biomarkers for disease progression and response to treatment. Such co-expressed genes can be assayed in lieu of, or in addition to, assaying of the prognostic and/or predictive gene with which they are co-expressed.

[00123] One skilled in the art will recognize that many co-expression analysis methods now known or later developed will fall within the scope and spirit of the present invention. These methods may incorporate, for example, correlation coefficients, co-expression network analysis, clique analysis, etc., and may be based on expression data from RT-PCR, microarrays, sequencing, and other similar technologies. For example, gene expression clusters can be identified using pair-wise analysis of correlation based on Pearson or Spearman correlation coefficients. (See, e.g., Pearson K. and Lee A., *Biometrika* 2, 357 (1902); C. Spearman, *Amer. J. Psychol* 15:72-101 (1904); J. Myers, A. Well, *Research Design and Statistical Analysis*, p. 508 (2nd Ed., 2003).) In general, a correlation coefficient of equal to or greater than 0.3 is considered to be statistically significant in a sample size of at least 20. (See, e.g., G. Norman, D. Streiner, *Biostatistics: The Bare Essentials*, 137-138 (3rd Ed. 2007).)

[00124] All aspects of the present disclosure may also be practiced such that a limited number of additional genes that are co-expressed with the disclosed genes, for example as

evidenced by high Pearson correlation coefficients, are included in a prognostic or predictive test in addition to and/or in place of disclosed genes.

[00125] Having described the invention, the same will be more readily understood through reference to the following Examples, which are provided by way of illustration, and are not intended to limit the invention in any way. All citations throughout the disclosure are hereby expressly incorporated by reference.

EXAMPLES

[00126] Two studies were performed to demonstrate the feasibility of gene expression profiling from renal tumors obtained from renal cell carcinoma patients. (See Abstract by M. Zhou, et al., Optimized RNA extraction and RT-PCR assays provide successful molecular analysis on a wide variety of archival fixed tissues, AACR Annual Meeting (2007)).

Study Design

[00127] Renal tumor tissue was obtained from approximately 1200 patients from the Cleveland Clinic Foundation (CCF) database. This database consists of patients who were diagnosed with renal carcinoma, clear cell type, stage I, II and III between the years of 1985 and 2003, who had available paraffin-embedded tumor (PET) blocks and adequate clinical follow-up, and who were not treated with adjuvant/neo-adjuvant systemic therapy. Patients with inherited VHL disease or bilateral tumors were also excluded. Tumors were graded using (1) Fuhrman grading system as noted in the World Health Organization Classification of Tumours: Pathology and Genetics: Tumours of the Urinary System and Male Genital Organs; and (2) the modified Fuhrman grading system (Table 1). In general, if no nodal involvement is expected or observed for patients, inspection of nodal involvement is not conducted and Nx is noted. In this study, Nx was treated as N0 for purposes of stage classification. The expression of 732 genes was quantitatively assessed for each patient tissue sample.

Table 1: Fuhrman Grading Systems

	Fuhrman Grade (Modified)	Fuhrman Grade (WHO)
1	Nuclei small as lymphocyte with condensed chromatin	Small, round, uniform nuclei (~ 10 um); nucleoli absent or inconspicuous (at 400x)
2	Nuclei both small as lymphocytes with condensed chromatin and other nuclei demonstrating enlarged, open chromatin	Larger nuclei (~ 15um) with irregular outline; small nucleoli present (at 400x)
3	All nuclei enlarged with open chromatin	Larger nuclei (approaching 20um) with more irregular outline; prominent nucleoli present (at 100x)
4	Large bizarre nuclei	Grade 3 features with pleomorphic or multilobed nuclei, with or without spindle cells

Inclusion Criteria

[00128] (1) Patients who underwent nephrectomy at CCF and who have a minimum of 6 months clinical follow-up or have recurrent RCC, documented in the clinical chart, database or registry.

[00129] (2) Diagnosed with RCC, clear cell type, stage I, II, or III.

[00130] (3) Renal blocks fixed in Formalin, Hollandes fluid or Zenkers fixative.

Exclusion Criteria

[00131] (1) No tumor block available from initial diagnosis in the Cleveland Clinic archive.

[00132] (2) No tumor or very little tumor (<5% of invasive cancer cell area) in block as assessed by examination of the Hollandes and/or hematoxylin and eosin stained (H&E) slide.

[00133] (3) High cycle threshold (C_t) values of reference genes. All samples regardless of their RNA amount will be tested by RT-qPCR, but only plates where the average C_t of reference genes is less than 35 will be analyzed.

[00134] (4) Patients with inherited VHL disease and/or bilateral tumors

[00135] (5) Patients who received neo-adjuvant or adjuvant systemic therapy

Concordance for clinical and pathologic factors

[00136] Two separate pathology laboratories conducted analyses of several clinical and pathologic factors using the same standardized measures. The level of concordance, by

covariate, is provided in Table 2, below. For purposes of the statistical analysis, the determination of only one of the central laboratories was used.

Table 2: Concordance between two central laboratories for clinical/pathologic covariates

Covariate	Method of analysis	Concordance
Presence of necrosis	Microscopic technique per Leibovich BC et al. (2003) Cancer 97(3), 1663-1671.	47%
Tumor grade	Fuhrman	65%

Expression Profile Gene Panel

[00137] The RNA from paraffin embedded tissue (PET) samples obtained from 942 patients who met all inclusion/exclusion criteria was extracted using protocols optimized for fixed renal tissue and perform molecular assays of quantitative gene expression using TaqMan® RT-PCR. RT-PCR was performed with RNA input at 1.0 ng per 5 µL-reaction volume using two 384 well plates.

[00138] RT-PCR analysis of PET samples was conducted using 732 genes. These genes were evaluated for association with the primary and secondary endpoints, recurrence-free interval (RFI), disease-free survival (DFS) and overall survival (OS).

[00139] All primary and secondary analyses were conducted on reference normalized gene expression levels using the mean of the reference genes for normalization. Three normalization schemes were tested using AAMP, ARF1, ATP5E, EEF1A1, GPX1, RPS23, SDHA, UBB, and RPLP1. Some or all of the other genes in the test panel were used in analyses of alternative normalization schemes.

[00140] Of the 732 genes, 647 were deemed evaluable for further consideration. The outcome analyses of the 647 evaluable genes were adjusted for multiple hypothesis tests, by allowing for a 10% false discovery rate (FDR), using Storey's procedure, and using True Discovery Rate Degree of Association Sets (TDRDAS) with separate classes (M. Crager, Gene identification using true discovery rate degree of association sets and estimates corrected for regression to the mean, *Statistics in Medicine* 29:33-45 (2009)). Unadjusted for baseline covariates and without controlling the FDR, a subset of 448 (69%) of genes were identified as

significantly associated with RFI ($p \leq 0.05$). Additional analysis was conducted using a supervised principal component analysis (PCA) on a subset of 188 genes that have maximum lower bound (MLB) > 1.2 using TDRDAS analysis, controlling the FDR but not taking into account the separate classes. The top 10 factors were modified by keeping the genes with high factor loadings (loading/max loading > 0.7) were kept. The modified top 10 factors were put into a 5-fold cross validation to assess performance of the gene groups and identify factors that were appearing most frequently.

[00141] Genes that had a significant association ($p \leq 0.05$) with risk of recurrence are listed in Tables 3a and 3b, wherein genes that are positively associated with a good prognosis (i.e., increased expression indicates a lower risk of recurrence) are listed in Table 3a. Genes that are negatively associated with a good prognosis (i.e., increased expression indicates a higher risk of recurrence) are listed in Table 3b. Genes that were associated, positively or negatively, with a good prognosis but not associated with clinical/pathologic covariates are BBC3, CCR7, CCR4, and VCAN. In addition, genes associated positively with a good prognosis after adjusting for clinical and pathologic covariates (stage, tumor grade, tumor size, nodal status, and presence of necrosis) are listed in Table 8a. Genes associated negatively with a good prognosis after adjusting for clinical/pathologic covariates are listed in Table 8b. Of those genes listed in Tables 8a and 8b, 16 genes with significant association, positively or negatively, with a good prognosis after adjusting for clinical and pathologic covariates and controlling the false discovery rate at 10% are listed in Table 9.

[00142] For the majority of these genes significantly associated with RFI ($p \leq 0.05$) (82%), increased expression is associated with a good prognosis. Most of the genes significantly associated with RFI showed consistency between (1) between stages (I-III); (2) primary and secondary endpoints (RFI and OS). See, e.g., Figures 1 and 2, respectively.

[00143] From this analysis, certain gene subsets emerged as significantly associated with recurrence and overall survival. For example, increased expression of angiogenesis genes (e.g., EMCN, PPAP2B, NOS3, NUDT6, PTPRB, SNRK, APOLD1, PRKCH, and CEACAM1), cell adhesion/extracellular matrix genes (e.g., ITGB5, ITGB1, A2M, TIMP3), immune response genes (e.g., CCL5, CCXL9, CCR7, IL8, IL6, and CX3CL1), cell cycle (e.g., BUB1, TPX2), apoptosis (e.g., CASP10), and transport genes (AQP1) were strongly associated, positively or negatively, with RFI.

[00144] Also, certain genes that are associated with pathway targets for renal cancer drugs (sunitinib, sorafenib, temsirolimus, bevacizumab, everolimus, pazopanib) were identified as having a significant association with outcome, including: KIT, PDGFA, PDGFB, PDGFC, PDGFD, PDGFRb, KRAS, RAF1, MTOR, HIF1AN, VEGFA, VEGFB, and FLT4.

[00145] It was determined that the presence of necrosis in these tumors was associated with a higher risk of recurrence, at least in the first 4 years after surgery. See Figure 3. However, the prognostic effect of necrosis after year 4 was negligible.

[00146] It was also determined that expression of certain genes was correlated, positively or negatively, with pathologic and/or clinical factors (“proxy genes”). For example, increased expression of the proxy genes listed in Tables 4a-7b correlate, positively or negatively, with tumor stage, tumor grade, presence of necrosis, and nodal invasion, respectively. In Tables 4a-7b, gene expression was normalized and then standardized such that the odds ratio (OR) reflects a one standard deviation change in gene expression.

[00147] From these, key genes were identified as good proxies for baseline covariates (stage, grade, necrosis), including TSPAN7, TEK, LDB2, TIMP3, SHANK3, RGS5, KDR, SDPR, EPAS1, ID1, TGFBR2, FLT4, SDPR, ENDRB, JAG1, DLC1, and KL. Several of these genes are in the hypoxia-induced pathway: SHANK3, RGS5, EPAS1, KDR, JAG1, TGFBR2, FLT4, SDPR, DLC1, EDNRB.

[00148] Figures 4 – 7 provide a comparison of patient stratification (Low, Intermediate, or High Risk) obtained by applying the Mayo prognostic tool (described in Leibovich et al. “prediction of progression after radical nephrectomy for patients with clear cell renal cell carcinoma” (2003) Cancer 97:1663-1671) CCF expression data. As shown in Figure 4, the Mayo prognostic tool alone provides for stratification into three populations Low Risk (93% recurrence free at 5 years), Intermediate Risk (79% recurrence free at 5 years), and High Risk (36% recurrence free at 5 years). In contrast, use of expression data from even one gene (as exemplified by EMCN (Fig. 5.), AQP1 (Fig. 6), or PPAP2B (Fig. 7)) allowed for more detailed stratification of patients according to risk.

Table 3a: Genes for which increased expression is associated with lower risk of cancer recurrence (p-value $\leq .05$)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
YB-1.2	YBX1	<0.0001	0.75	IGF1R.3	IGF1R	<0.0001	0.58
XIAP.1	XIAP	0.0009	0.81	IFI27.1	IFI27	0.0205	0.85
WWOX.5	WWOX	<0.0001	0.71	ID3.1	ID3	<0.0001	0.69
VWF.1	VWF	<0.0001	0.61	ID2.4	ID2	<0.0001	0.75
VEGF.1	VEGFA	<0.0001	0.75	ID1.1	ID1	<0.0001	0.55
VCAM1.1	VCAM1	<0.0001	0.66	ICAM2.1	ICAM2	<0.0001	0.64
USP34.1	USP34	0.0003	0.79	HYAL2.1	HYAL2	<0.0001	0.60
UMOD.1	UMOD	<0.0001	0.68	HYAL1.1	HYAL1	<0.0001	0.58
UGCG.1	UGCG	<0.0001	0.71	HSPG2.1	HSPG2	<0.0001	0.60
UBB.1	UBB	<0.0001	0.62	HSD11B2.1	HSD11B2	<0.0001	0.63
UBE1C.1	UBA3	0.0003	0.79	Hepsin.1	HPN	0.0001	0.76
TS.1	TYMS	0.0056	0.83	HPCAL1.1	HPCAL1	0.0031	0.82
tusc4.2	TUSC4	<0.0001	0.76	HMGB1.1	HMGB1	<0.0001	0.67
TSPAN7.2	TSPAN7	<0.0001	0.52	HLA-DPB1.1	HLA-DPB1	<0.0001	0.70
TSC2.1	TSC2	0.0043	0.82	HIF1AN.1	HIF1AN	<0.0001	0.77
TSC1.1	TSC1	<0.0001	0.71	HDAC1.1	HDAC1	0.0003	0.78
P53.2	TP53	0.0008	0.81	HAVCR1.1	HAVCR1	0.0003	0.79
TOP2B.2	TOP2B	0.0001	0.80	HADH.1	HADH	<0.0001	0.68
TNFSF12.1	TNFSF12	<0.0001	0.68	GZMA.1	GZMA	0.0125	0.84
TRAIL.1	TNFSF10	0.0065	0.83	GSTp.3	GSTP1	<0.0001	0.76
TNFRSF11B.1	TNFRSF11B	0.0002	0.78	GSTM3.2	GSTM3	<0.0001	0.70
TNFRSF10D.1	TNFRSF10D	<0.0001	0.76	GSTM1.1	GSTM1	<0.0001	0.72
DR5.2	TNFRSF10B	<0.0001	0.75	GRB7.2	GRB7	<0.0001	0.74
TNFAIP6.1	TNFAIP6	0.0005	0.79	GPX3.1	GPX3	<0.0001	0.75
TMEM47.1	TMEM47	<0.0001	0.57	GJA1.1	GJA1	0.0049	0.84
TMEM27.1	TMEM27	<0.0001	0.58	GFRA1.1	GFRA1	0.0003	0.77
TLR3.1	TLR3	<0.0001	0.72	GCLC.3	GCLC	<0.0001	0.68
TIMP3.3	TIMP3	<0.0001	0.50	GBP2.2	GBP2	0.0156	0.85
TIMP2.1	TIMP2	<0.0001	0.73	GATM.1	GATM	<0.0001	0.66
THBS1.1	THBS1	0.0002	0.79	GATA3.3	GATA3	0.0001	0.75
TGFBR2.3	TGFBR2	<0.0001	0.54	FOS.1	FOS	<0.0001	0.74

Table 3a: Genes for which increased expression is associated with lower risk of cancer recurrence (p-value $\leq$.05)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
TGFBR1.1	TGFBR1	<0.0001	0.71	FOLR1.1	FOLR1	0.0003	0.79
TGFB2.2	TGFB2	<0.0001	0.65	FLT4.1	FLT4	<0.0001	0.54
TGFA.2	TGFA	<0.0001	0.70	FLT3LG.1	FLT3LG	0.0001	0.73
TEK.1	TEK	<0.0001	0.47	FLT1.1	FLT1	<0.0001	0.59
TCF4.1	TCF4	<0.0001	0.62	FILIP1.1	FILIP1	<0.0001	0.72
TAP1.1	TAP1	0.0024	0.82	FIGF.1	FIGF	0.0014	0.76
TAGLN.1	TAGLN	<0.0001	0.70	FHL1.1	FHL1	<0.0001	0.68
TACSTD2.1	TACSTD2	<0.0001	0.72	FHIT.1	FHIT	<0.0001	0.73
SUCLG1.1	SUCLG1	<0.0001	0.72	FH.1	FH	<0.0001	0.73
STK11.1	STK11	<0.0001	0.76	FGFR2 isoform 1.1	FGFR2	<0.0001	0.70
STAT5B.2	STAT5B	<0.0001	0.71	FGFR1.3	FGFR1	0.0001	0.77
STAT5A.1	STAT5A	<0.0001	0.72	FGF2.2	FGF2	<0.0001	0.73
STAT3.1	STAT3	0.0016	0.80	FGF1.1	FGF1	0.0015	0.78
SPRY1.1	SPRY1	<0.0001	0.68	FDPS.1	FDPS	<0.0001	0.69
SPARCL1.1	SPARCL1	<0.0001	0.71	FBXW7.1	FBXW7	0.0001	0.75
SPARC.1	SPARC	0.0009	0.80	fas.1	FAS	<0.0001	0.73
SOD1.1	SOD1	0.0114	0.84	FABP1.1	FABP1	0.0455	0.86
SNRK.1	SNRK	<0.0001	0.54	ESRRG.3	ESRRG	0.0008	0.79
SNAI1.1	SNAI1	0.0055	0.82	ERG.1	ERG	<0.0001	0.60
MADH4.1	SMAD4	<0.0001	0.60	ERCC1.2	ERCC1	<0.0001	0.78
MADH2.1	SMAD2	<0.0001	0.64	ErbB3.1	ERBB3	0.0001	0.77
SLC34A1.1	SLC34A1	0.0004	0.76	HER2.3	ERBB2	<0.0001	0.65
SLC22A6.1	SLC22A6	0.0003	0.77	EPHB4.1	EPHB4	<0.0001	0.64
SKIL.1	SKIL	0.0004	0.79	EPHA2.1	EPHA2	<0.0001	0.58
SHANK3.1	SHANK3	<0.0001	0.53	EPAS1.1	EPAS1	<0.0001	0.55
SGK.1	SGK1	<0.0001	0.67	ENPP2.1	ENPP2	0.0090	0.84
FRP1.3	SFRP1	0.0355	0.86	ENPEP.1	ENPEP	<0.0001	0.74
SEMA3F.3	SEMA3F	<0.0001	0.67	CD105.1	ENG	<0.0001	0.60
SELENBP1.1	SELENBP1	0.0016	0.81	EMP1.1	EMP1	<0.0001	0.71
SDPR.1	SDPR	<0.0001	0.51	EMCN.1	EMCN	<0.0001	0.43
SDHA.1	SDHA	0.0003	0.77	ELTD1.1	ELTD1	<0.0001	0.76
SCNN1A.2	SCNN1A	0.0134	0.83	EIF2C1.1	EIF2C1	<0.0001	0.67

Table 3a: Genes for which increased expression is associated with lower risk of cancer recurrence (p-value $\leq .05$)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
SCN4B.1	SCN4B	<0.0001	0.68	EGR1.1	EGR1	<0.0001	0.72
KIAA1303 raptor.1	RPTOR	<0.0001	0.77	EGLN3.1	EGLN3	0.0002	0.80
RPS6KB1.3	RPS6KB1	<0.0001	0.73	EGFR.2	EGFR	0.0005	0.80
RPS6KAI.1	RPS6KA1	0.0291	0.86	EFNB2.1	EFNB2	<0.0001	0.64
RPS23.1	RPS23	<0.0001	0.73	EFNB1.2	EFNB1	<0.0001	0.68
ROCK2.1	ROCK2	<0.0001	0.66	EEF1A1.1	EEF1A1	<0.0001	0.64
ROCK1.1	ROCK1	<0.0001	0.58	EDNRB.1	EDNRB	<0.0001	0.56
RIPK1.1	RIPK1	<0.0001	0.64	EDN2.1	EDN2	0.0005	0.77
rhoC.1	RHOC	0.0213	0.86	EDN1 endothelin.1	EDN1	<0.0001	0.62
RhoB.1	RHOB	<0.0001	0.63	EBAG9.1	EBAG9	0.0041	0.82
ARHA.1	RHOA	<0.0001	0.64	DUSP1.1	DUSP1	0.0029	0.82
RGS5.1	RGS5	<0.0001	0.52	DPYS.1	DPYS	0.0112	0.84
FLJ22655.1	RERGL	<0.0001	0.51	DPEP1.1	DPEP1	<0.0001	0.64
NFKBp65.3	RELA	0.0027	0.82	DLL4.1	DLL4	<0.0001	0.76
RB1.1	RB1	<0.0001	0.73	DLC1.1	DLC1	<0.0001	0.55
RASSF1.1	RASSF1	0.0040	0.83	DKFZP564O08 23.1	DKFZP564O 0823	<0.0001	0.62
RARB.2	RARB	<0.0001	0.57	DICER1.2	DICER1	<0.0001	0.71
RALBP1.1	RALBP1	<0.0001	0.73	DIAPH1.1	DIAPH1	0.0009	0.80
RAF1.3	RAF1	0.0008	0.81	DIABLO.1	DIABLO	0.0134	0.84
PTPRG.1	PTPRG	<0.0001	0.57	DHPS.3	DHPS	<0.0001	0.70
PTPRB.1	PTPRB	<0.0001	0.51	DET1.1	DET1	<0.0001	0.74
PTN.1	PTN	<0.0001	0.71	DEFB1.1	DEFB1	0.0025	0.81
PTK2.1	PTK2	<0.0001	0.61	DDC.1	DDC	<0.0001	0.68
PTHR1.1	PTH1R	<0.0001	0.55	DCXR.1	DCXR	0.0081	0.83
PTEN.2	PTEN	<0.0001	0.74	DAPK1.3	DAPK1	<0.0001	0.60
PSMB9.1	PSMB9	0.0025	0.82	CYR61.1	CYR61	<0.0001	0.70
PSMB8.1	PSMB8	0.0239	0.85	CYP3A4.2	CYP3A4	0.0380	0.86
PRSS8.1	PRSS8	<0.0001	0.71	CXCL9.1	CXCL9	0.0362	0.87
PRPS2.1	PRPS2	0.0156	0.85	CXCL12.1	CXCL12	<0.0001	0.69
PRKCH.1	PRKCH	<0.0001	0.63	CX3CR1.1	CX3CR1	<0.0001	0.72
PPP2CA.1	PPP2CA	<0.0001	0.77	CX3CL1.1	CX3CL1	<0.0001	0.58

Table 3a: Genes for which increased expression is associated with lower risk of cancer recurrence (p-value $\leq .05$)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
PPARG.3	PPARG	<0.0001	0.59	CUL1.1	CUL1	0.0003	0.80
PPAP2B.1	PPAP2B	<0.0001	0.50	CUBN.1	CUBN	<0.0001	0.61
PLG.1	PLG	<0.0001	0.70	CTSS.1	CTSS	0.0016	0.82
PLAT.1	PLAT	<0.0001	0.64	CTSH.2	CTSH	<0.0001	0.78
PLA2G4C.1	PLA2G4C	<0.0001	0.67	B-Catenin.3	CTNNB1	<0.0001	0.74
PIK3CA.1	PIK3CA	<0.0001	0.75	A-Catenin.2	CTNNA1	<0.0001	0.77
PI3K.2	PIK3C2B	<0.0001	0.56	CTGF.1	CTGF	0.0004	0.79
PFKP.1	PFKP	0.0067	0.84	CSF1R.2	CSF1R	0.0015	0.82
CD31.3	PECAM1	<0.0001	0.60	CSF1.1	CSF1	0.0016	0.81
PDZK3.1	PDZK3	0.0005	0.77	CRADD.1	CRADD	0.0021	0.81
PDZK1.1	PDZK1	<0.0001	0.69	COL4A2.1	COL4A2	0.0016	0.80
PDGFRb.3	PDGFRB	<0.0001	0.74	COL18A1.1	COL18A1	0.0007	0.79
PDGFD.2	PDGFD	<0.0001	0.58	CLU.3	CLU	0.0151	0.85
PDGFC.3	PDGFC	<0.0001	0.66	CLDN7.2	CLDN7	0.0023	0.81
PDGFB.3	PDGFB	<0.0001	0.63	CLDN10.1	CLDN10	<0.0001	0.69
PDGFA.3	PDGFA	<0.0001	0.75	CLCNKB.1	CLCNKB	0.0002	0.74
PCK1.1	PCK1	<0.0001	0.65	CFLAR.1	CFLAR	<0.0001	0.65
PCCA.1	PCCA	<0.0001	0.66	CEACAM1.1	CEACAM1	<0.0001	0.59
PARD6A.1	PARD6A	0.0001	0.76	p27.3	CDKN1B	0.0018	0.83
Pak1.2	PAK1	0.0011	0.81	p21.3	CDKN1A	0.0027	0.81
PAH.1	PAH	0.0296	0.86	CDH6.1	CDH6	<0.0001	0.75
OGG1.1	OGG1	0.0051	0.84	CDH5.1	CDH5	<0.0001	0.59
BFGF.3	NUDT6	<0.0001	0.54	CDH16.1	CDH16	<0.0001	0.75
NRG1.3	NRG1	0.0049	0.81	CDH13.1	CDH13	<0.0001	0.66
NPR1.1	NPR1	<0.0001	0.73	CD4.1	CD4	0.0009	0.81
NPM1.2	NPM1	<0.0001	0.73	CD36.1	CD36	<0.0001	0.65
NOTCH3.1	NOTCH3	<0.0001	0.70	CD34.1	CD34	<0.0001	0.62
NOTCH2.1	NOTCH2	0.0040	0.84	CCR7.1	CCR7	0.0271	0.86
NOTCH1.1	NOTCH1	<0.0001	0.64	CCR4.2	CCR4	0.0106	0.83
NOS3.1	NOS3	<0.0001	0.55	CCND1.3	CCND1	<0.0001	0.70
NOS2A.3	NOS2	<0.0001	0.66	CCL4.2	CCL4	0.0012	0.80
NOL3.1	NOL3	0.0132	0.85	MCP1.1	CCL2	<0.0001	0.75
NFX1.1	NFX1	<0.0001	0.67	CAT.1	CAT	<0.0001	0.72

Table 3a: Genes for which increased expression is associated with lower risk of cancer recurrence (p-value $\leq$.05)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
NFKBp50.3	NFKB1	0.0001	0.77	CASP6.1	CASP6	0.0369	0.87
NFATC2.1	NFATC2	0.0003	0.78	CASP10.1	CASP10	<0.0001	0.69
NFAT5.1	NFAT5	0.0010	0.82	CALD1.2	CALD1	<0.0001	0.61
MYRIP.2	MYRIP	0.0002	0.75	CA9.3	CA9	0.0035	0.84
MYH11.1	MYH11	<0.0001	0.61	CA2.1	CA2	0.0006	0.79
cMYC.3	MYC	0.0002	0.79	C7.1	C7	0.0030	0.82
MVP.1	MVP	0.0052	0.83	ECRG4.1	C2orf40	<0.0001	0.57
MUC1.2	MUC1	0.0405	0.87	C13orf15.1	C13orf15	<0.0001	0.57
FRAP1.1	MTOR	<0.0001	0.75	BUB3.1	BUB3	0.0002	0.77
MSH3.2	MSH3	<0.0001	0.75	BTRC.1	BTRC	0.0006	0.81
MSH2.3	MSH2	<0.0001	0.71	CIAP1.2	BIRC2	0.0030	0.82
GBL.1	MLST8	0.0246	0.87	BIN1.3	BIN1	0.0005	0.80
MIF.2	MIF	0.0012	0.80	BGN.1	BGN	<0.0001	0.76
MICA.1	MICA	<0.0001	0.71	BCL2L12.1	BCL2L12	0.0322	0.86
MGMT.1	MGMT	<0.0001	0.68	Bclx.2	BCL2L1	<0.0001	0.74
MCM3.3	MCM3	0.0188	0.85	Bcl2.2	BCL2	<0.0001	0.57
MCAM.1	MCAM	<0.0001	0.71	BBC3.2	BBC3	0.0449	0.87
MARCKS.1	MARCKS	0.0301	0.87	BAG1.2	BAG1	<0.0001	0.64
ERK1.3	MAPK3	<0.0001	0.65	BAD.1	BAD	0.0076	0.85
ERK2.3	MAPK1	0.0005	0.79	ATP6V1B1.1	ATP6V1B1	0.0001	0.71
MAP4.1	MAP4	<0.0001	0.65	ASS1.1	ASS1	<0.0001	0.75
MAP2K3.1	MAP2K3	<0.0001	0.69	ARRB1.1	ARRB1	<0.0001	0.62
MAP2K1.1	MAP2K1	0.0046	0.83	ARHGDIB.1	ARHGDIB	<0.0001	0.66
MAL2.1	MAL2	0.0001	0.76	AQP1.1	AQP1	<0.0001	0.50
MAL.1	MAL	<0.0001	0.66	APOLD1.1	APOLD1	<0.0001	0.57
LYZ.1	LYZ	0.0458	0.88	APC.4	APC	<0.0001	0.73
LTF.1	LTF	0.0005	0.76	ANXA4.1	ANXA4	0.0018	0.81
LRP2.1	LRP2	<0.0001	0.67	ANXA1.2	ANXA1	0.0009	0.80
LMO2.1	LMO2	<0.0001	0.74	ANTXR1.1	ANTXR1	0.0043	0.82
LDB2.1	LDB2	<0.0001	0.52	ANGPTL4.1	ANGPTL4	0.0033	0.84
LDB1.2	LDB1	<0.0001	0.71	ANGPTL3.3	ANGPTL3	0.0003	0.75
LAMA4.1	LAMA4	0.0279	0.86	ANGPT1.1	ANGPT1	<0.0001	0.57
KRT7.1	KRT7	<0.0001	0.68	ALDOB.1	ALDOB	<0.0001	0.62

Table 3a: Genes for which increased expression is associated with lower risk of cancer recurrence (p-value $\leq$.05)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
K-ras.10	KRAS	<0.0001	0.72	ALDH6A1.1	ALDH6A1	<0.0001	0.63
KL.1	KL	<0.0001	0.55	ALDH4.2	ALDH4A1	0.0172	0.85
Kitlng.4	KITLG	<0.0001	0.67	AKT3.2	AKT3	<0.0001	0.57
c-kit.2	KIT	<0.0001	0.72	AKT2.3	AKT2	<0.0001	0.75
KDR.6	KDR	<0.0001	0.54	AKT1.3	AKT1	<0.0001	0.71
KCNJ15.1	KCNJ15	<0.0001	0.63	AIF1.1	AIF1	0.0349	0.87
HTATIP.1	KAT5	<0.0001	0.64	AHR.1	AHR	<0.0001	0.74
G-Catenin.1	JUP	<0.0001	0.64	AGTR1.1	AGTR1	<0.0001	0.57
AP-1 (JUN official).2	JUN	0.0001	0.76	ADH1B.1	ADH1B	0.0002	0.77
JAG1.1	JAG1	<0.0001	0.55	ADFP.1	ADFP	0.0332	0.88
ITGB1.1	ITGB1	0.0085	0.83	ADD1.1	ADD1	<0.0001	0.58
ITGA7.1	ITGA7	<0.0001	0.66	ADAMTS5.1	ADAMTS5	0.0010	0.78
ITGA6.2	ITGA6	<0.0001	0.63	ADAMTS1.1	ADAMTS1	0.0056	0.83
ITGA4.2	ITGA4	<0.0001	0.74	ACE2.1	ACE2	<0.0001	0.62
ITGA3.2	ITGA3	0.0002	0.79	ACADSB.1	ACADSB	<0.0001	0.71
IQGAP2.1	IQGAP2	<0.0001	0.69	BCRP.1	ABCG2	<0.0001	0.58
INSR.1	INSR	<0.0001	0.67	MRP4.2	ABCC4	<0.0001	0.74
IMP3.1	IMP3	<0.0001	0.69	MRP3.1	ABCC3	0.0107	0.85
IL6ST.3	IL6ST	<0.0001	0.66	MRP1.1	ABCC1	<0.0001	0.75
IL15.1	IL15	<0.0001	0.70	ABCB1.5	ABCB1	0.0093	0.84
IGFBP6.1	IGFBP6	0.0004	0.79	NPD009 (ABAT official).3	ABAT	0.0001	0.76
IGFBP3.1	IGFBP3	0.0394	0.87	AAMP.1	AAMP	0.0008	0.80
IGFBP2.1	IGFBP2	0.0134	0.84	A2M.1	A2M	<0.0001	0.56

Table 3b: Genes for which increased expression is associated with higher risk of cancer recurrence (p-value $\leq .05$)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
WT1.1	WT1	0.0002	1.25	LGALS1.1	LGALS1	0.0017	1.25
VTN.1	VTN	0.0097	1.17	LAMB3.1	LAMB3	<0.0001	1.34
VDR.2	VDR	0.0031	1.22	LAMB1.1	LAMB1	0.0014	1.25
VCAN.1	VCAN	0.0036	1.22	L1CAM.1	L1CAM	0.0199	1.16
UBE2T.1	UBE2T	<0.0001	1.38	IL-8.1	IL8	<0.0001	1.53
C20 orf1.1	TPX2	<0.0001	1.76	IL6.3	IL6	<0.0001	1.41
TOP2A.4	TOP2A	<0.0001	1.39	ICAM1.1	ICAM1	0.0013	1.23
TK1.2	TK1	0.0018	1.22	HIST1H1D.1	HIST1H1D	0.0066	1.21
TIMP1.1	TIMP1	0.0259	1.16	FN1.1	FN1	0.0105	1.19
TGFBI.1	TGFBI	0.0004	1.26	F3.1	F3	<0.0001	1.31
SQSTM1.1	SQSTM1	0.0089	1.20	F2.1	F2	<0.0001	1.30
OPN, osteopontin.3	SPP1	<0.0001	1.43	ESPL1.3	ESPL1	0.0155	1.17
SPHK1.1	SPHK1	0.0025	1.22	EPHB2.1	EPHB2	0.0456	1.14
SLC7A5.2	SLC7A5	<0.0001	1.38	EPHB1.3	EPHB1	0.0007	1.22
SLC2A1.1	SLC2A1	0.0010	1.26	ENO2.1	ENO2	<0.0001	1.38
SLC16A3.1	SLC16A3	<0.0001	1.38	EIF4EBP1.1	EIF4EBP1	0.0098	1.19
SLC13A3.1	SLC13A3	0.0192	1.16	CXCR4.3	CXCR4	0.0066	1.21
SHC1.1	SHC1	0.0086	1.19	GRO1.2	CXCL1	<0.0001	1.30
SFN.1	SFN	0.0001	1.26	CTSB.1	CTSB	0.0233	1.17
SERPINA5.1	SERPINA5	0.0462	1.13	CRP.1	CRP	0.0314	1.13
SEMA3C.1	SEMA3C	<0.0001	1.45	CP.1	CP	0.0002	1.32
SAA2.2	SAA2	<0.0001	1.59	COL7A1.1	COL7A1	0.0003	1.24
S100A1.1	S100A1	0.0348	1.16	COL1A1.1	COL1A1	0.0029	1.23
RRM2.1	RRM2	0.0002	1.27	Chk1.2	CHEK1	0.0002	1.26
RPLP1.1	RPLP1	0.0049	1.22	CENPF.1	CENPF	<0.0001	1.36
PTTG1.2	PTTG1	<0.0001	1.45	CD82.3	CD82	0.0009	1.25
COX2.1	PTGS2	0.0013	1.22	CD44s.1	CD44_s	0.0065	1.21
PLAUR.3	PLAUR	<0.0001	1.33	CCNE1.1	CCNE1	0.0098	1.17
PF4.1	PF4	0.0034	1.20	CCNB1.2	CCNB1	<0.0001	1.42
PCSK6.1	PCSK6	0.0269	1.17	CCL20.1	CCL20	0.0029	1.22
MYBL2.1	MYBL2	<0.0001	1.33	CA12.1	CA12	<0.0001	1.48
MT1X.1	MT1X	0.0070	1.20	C3.1	C3	0.0176	1.18
MMP9.1	MMP9	<0.0001	1.54	BUB1.1	BUB1	<0.0001	1.59

Table 3b: Genes for which increased expression is associated with higher risk of cancer recurrence (p-value $\leq$.05)

Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI		Gene	Official Symbol	Univariate Cox Analyses (No Covariate Adjustment) with RFI	
		p-value for HR	HR			p-value for HR	HR
MMP7.1	MMP7	0.0312	1.15	SURV.2	BIRC5	<0.0001	1.37
MMP14.1	MMP14	<0.0001	1.47	ciAP2.2	BIRC3	0.0484	1.15
Ki-67.2	MKI67	<0.0001	1.33	BCL2A1.1	BCL2A1	0.0483	1.11
mGST1.2	MGST1	<0.0001	1.38	STK15.2	AURKA	0.0002	1.28
MDK.1	MDK	0.0001	1.31	ANXA2.2	ANXA2	0.0315	1.16
LOX.1	LOX	<0.0001	1.42	ALOX5.1	ALOX5	0.0473	1.14
LMNB1.1	LMNB1	<0.0001	1.40	ADAM8.1	ADAM8	0.0002	1.29
LIMK1.1	LIMK1	<0.0001	1.43	MRP2.3	ABCC2	0.0004	1.28

Table 4a: Proxy genes for which increased expression is associated with higher tumor stage (p-value $\leq$.05)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
WT1.1	WT1	<0.0001	1.41	LAMA3.1	LAMA3	0.0121	1.22
VTN.1	VTN	0.0007	1.29	L1CAM.1	L1CAM	0.0091	1.22
VDR.2	VDR	0.0065	1.25	ISG20.1	ISG20	0.0006	1.34
UBE2T.1	UBE2T	<0.0001	1.61	IL-8.1	IL8	<0.0001	1.89
TSPAN8.1	TSPAN8	0.0072	1.23	IL6.3	IL6	<0.0001	1.68
C20 orf1.1	TPX2	<0.0001	1.89	IGF1.2	IGF1	0.0214	1.20
TOP2A.4	TOP2A	<0.0001	1.55	ICAM1.1	ICAM1	<0.0001	1.42
TK1.2	TK1	0.0001	1.34	HIST1H1D.1	HIST1H1D	0.0005	1.33
TIMP1.1	TIMP1	0.0021	1.29	GPX2.2	GPX2	0.0129	1.22
TGFB1.1	TGFB1	0.0001	1.39	FN1.1	FN1	0.0002	1.36
OPN, osteopontin.3	SPP1	0.0001	1.38	FAP.1	FAP	0.0455	1.18
SLC7A5.2	SLC7A5	<0.0001	1.51	F3.1	F3	<0.0001	1.52
SLC2A1.1	SLC2A1	0.0081	1.24	F2.1	F2	<0.0001	1.79
SLC16A3.1	SLC16A3	<0.0001	1.46	ESPL1.3	ESPL1	0.0001	1.35
SFN.1	SFN	0.0001	1.36	EPB41L3.1	EPB41L3	0.0067	1.24
SEMA3C.1	SEMA3C	<0.0001	1.42	ENO2.1	ENO2	0.0016	1.31
SELL.1	SELL	0.0313	1.19	EIF4EBP1.1	EIF4EBP1	0.0036	1.27
SAA2.2	SAA2	<0.0001	2.04	E2F1.3	E2F1	0.0017	1.27
RRM2.1	RRM2	<0.0001	1.47	DCN.1	DCN	0.0152	1.22
RPLP1.1	RPLP1	0.0007	1.33	CXCR6.1	CXCR6	0.0013	1.30
RAD51.1	RAD51	0.0010	1.31	BLR1.1	CXCR5	0.0232	1.19
PTTG1.2	PTTG1	<0.0001	1.61	CXCR4.3	CXCR4	0.0003	1.35
COX2.1	PTGS2	0.0011	1.29	GRO1.2	CXCL1	0.0005	1.31
PTGIS.1	PTGIS	0.0034	1.27	CTSB.1	CTSB	0.0110	1.24
PLAUR.3	PLAUR	<0.0001	1.51	CRP.1	CRP	0.0002	1.31
PF4.1	PF4	0.0027	1.26	CP.1	CP	0.0008	1.34
PDGFRa.2	PDGFRA	0.0480	1.17	COL7A1.1	COL7A1	0.0010	1.28
PCSK6.1	PCSK6	0.0041	1.27	COL1A1.1	COL1A1	0.0001	1.40
NNMT.1	NNMT	0.0003	1.34	Chk2.3	CHEK2	0.0050	1.27
NME2.1	NME2	0.0028	1.28	Chk1.2	CHEK1	<0.0001	1.43
MYBL2.1	MYBL2	<0.0001	1.50	CENPF.1	CENPF	<0.0001	1.55
MT1X.1	MT1X	0.0192	1.21	CD82.3	CD82	0.0001	1.38
MMP9.1	MMP9	<0.0001	1.79	CD44s.1	CD44_s	0.0060	1.25
MMP7.1	MMP7	0.0252	1.20	CCNE2.2	CCNE2_2	0.0229	1.19
MMP14.1	MMP14	<0.0001	1.88	CCNB1.2	CCNB1	<0.0001	1.60

Table 4a: Proxy genes for which increased expression is associated with higher tumor stage (p-value $\leq$.05)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
Ki-67.2	MKI67	<0.0001	1.48	CCL20.1	CCL20	0.0010	1.30
mGST1.2	MGST1	0.0004	1.37	CA12.1	CA12	<0.0001	1.66
MDK.1	MDK	<0.0001	1.42	C3.1	C3	0.0009	1.32
MDH2.1	MDH2	0.0321	1.19	BUB1.1	BUB1	<0.0001	1.82
LRRC2.1	LRRC2	0.0259	1.19	SURV.2	BIRC5	<0.0001	1.46
LOX.1	LOX	<0.0001	1.78	BCL2A1.1	BCL2A1	<0.0001	1.44
LMNB1.1	LMNB1	<0.0001	1.82	STK15.2	AURKA	0.0002	1.36
LIMK1.1	LIMK1	<0.0001	1.46	APOL1.1	APOL1	0.0028	1.27
LAPTM5.1	LAPTM5	0.0102	1.23	ANXA2.2	ANXA2	0.0174	1.21
LAMB3.1	LAMB3	<0.0001	1.53	ADAM8.1	ADAM8	<0.0001	1.58
LAMB1.1	LAMB1	0.0452	1.18	MRP2.3	ABCC2	<0.0001	1.45

Table 4b: Proxy genes for which increased expression is associated with lower tumor stage (p-value $\leq .05$)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
YB-1.2	YBX1	<0.0001	0.63	IL-7.1	IL7	0.0444	0.85
XIAP.1	XIAP	<0.0001	0.62	IL6ST.3	IL6ST	<0.0001	0.50
WWOX.5	WWOX	0.0042	0.79	IL15.1	IL15	<0.0001	0.67
WISP1.1	WISP1	0.0096	0.81	IGFBP6.1	IGFBP6	0.0001	0.73
VWF.1	VWF	<0.0001	0.46	IGFBP3.1	IGFBP3	0.0191	0.83
VEGF.1	VEGFA	<0.0001	0.63	IGF1R.3	IGF1R	<0.0001	0.48
VCAM1.1	VCAM1	<0.0001	0.55	ID3.1	ID3	<0.0001	0.54
USP34.1	USP34	0.0001	0.72	ID2.4	ID2	0.0008	0.77
UMOD.1	UMOD	<0.0001	0.47	ID1.1	ID1	<0.0001	0.34
UGCG.1	UGCG	<0.0001	0.68	ICAM2.1	ICAM2	<0.0001	0.58
UBB.1	UBB	<0.0001	0.54	HYAL2.1	HYAL2	<0.0001	0.41
UBE1C.1	UBA3	<0.0001	0.62	HYAL1.1	HYAL1	<0.0001	0.44
TS.1	TYMS	0.0010	0.76	HSPG2.1	HSPG2	<0.0001	0.44
tusc4.2	TUSC4	<0.0001	0.57	HSD11B2.1	HSD11B2	<0.0001	0.47
TUSC2.1	TUSC2	0.0295	0.83	Hepsin.1	HPN	0.0031	0.79
TSPAN7.2	TSPAN7	<0.0001	0.35	HPCAL1.1	HPCAL1	0.0004	0.75
TSC2.1	TSC2	<0.0001	0.66	HNRPAB.3	HNRNPAB	0.0039	0.78
TSC1.1	TSC1	<0.0001	0.56	HMGB1.1	HMGB1	<0.0001	0.46
P53.2	TP53	<0.0001	0.67	HLA-DPB1.1	HLA-DPB1	<0.0001	0.55
TOP2B.2	TOP2B	<0.0001	0.69	HIF1AN.1	HIF1AN	<0.0001	0.57
TNIP2.1	TNIP2	0.0359	0.85	HIF1A.3	HIF1A	0.0076	0.80
TNFSF12.1	TNFSF12	<0.0001	0.50	HGF.4	HGF	0.0067	0.80
TRAIL.1	TNFSF10	0.0011	0.77	HDAC1.1	HDAC1	<0.0001	0.61
TNFRSF11B.1	TNFRSF11B	<0.0001	0.63	HAVCR1.1	HAVCR1	0.0001	0.73
TNFRSF10D.1	TNFRSF10D	<0.0001	0.57	HADH.1	HADH	<0.0001	0.62
DR5.2	TNFRSF10B	<0.0001	0.64	GSTT1.3	GSTT1	0.0112	0.82
TNFAIP6.1	TNFAIP6	0.0001	0.74	GSTp.3	GSTP1	<0.0001	0.64
TNF.1	TNF	0.0138	0.80	GSTM3.2	GSTM3	<0.0001	0.57
TMEM47.1	TMEM47	<0.0001	0.41	GSTM1.1	GSTM1	<0.0001	0.55
TMEM27.1	TMEM27	<0.0001	0.53	GRB7.2	GRB7	<0.0001	0.66
TLR3.1	TLR3	<0.0001	0.68	GRB14.1	GRB14	0.0123	0.81
TIMP3.3	TIMP3	<0.0001	0.39	GPX3.1	GPX3	0.0120	0.82
TIMP2.1	TIMP2	<0.0001	0.68	GNAS.1	GNAS	0.0003	0.74
THBS1.1	THBS1	<0.0001	0.65	GJA1.1	GJA1	0.0034	0.79
TGFBR2.3	TGFBR2	<0.0001	0.41	GFRA1.1	GFRA1	0.0164	0.82

Table 4b: Proxy genes for which increased expression is associated with lower tumor stage (p-value $\leq .05$)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
TGFBR1.1	TGFBR1	<0.0001	0.59	GCLM.2	GCLM	0.0056	0.80
TGFB2.2	TGFB2	<0.0001	0.50	GCLC.3	GCLC	<0.0001	0.49
TGFb1.1	TGFB1	<0.0001	0.67	GBP2.2	GBP2	0.0388	0.84
TGFA.2	TGFA	<0.0001	0.63	GATM.1	GATM	<0.0001	0.59
TEK.1	TEK	<0.0001	0.34	GATA3.3	GATA3	0.0002	0.72
TCF4.1	TCF4	<0.0001	0.47	FOS.1	FOS	<0.0001	0.65
TAP1.1	TAP1	0.0017	0.77	FOLR1.1	FOLR1	<0.0001	0.65
TAGLN.1	TAGLN	0.0001	0.72	FLT4.1	FLT4	<0.0001	0.37
TACSTD2.1	TACSTD2	<0.0001	0.58	FLT3LG.1	FLT3LG	<0.0001	0.61
SUCLG1.1	SUCLG1	<0.0001	0.56	FLT1.1	FLT1	<0.0001	0.40
STK11.1	STK11	<0.0001	0.58	FILIP1.1	FILIP1	<0.0001	0.47
STAT5B.2	STAT5B	<0.0001	0.50	FIGF.1	FIGF	<0.0001	0.53
STAT5A.1	STAT5A	<0.0001	0.60	FHL1.1	FHL1	<0.0001	0.52
STAT3.1	STAT3	0.0001	0.72	FHIT.1	FHIT	<0.0001	0.54
STAT1.3	STAT1	0.0344	0.84	FH.1	FH	<0.0001	0.67
SPRY1.1	SPRY1	<0.0001	0.57	FGFR2 isoform 1.1	FGFR2	<0.0001	0.62
SPAST.1	SPAST	0.0142	0.82	FGFR1.3	FGFR1	<0.0001	0.63
SPARCL1.1	SPARCL1	<0.0001	0.59	FGF2.2	FGF2	<0.0001	0.58
SPARC.1	SPARC	<0.0001	0.69	FGF1.1	FGF1	<0.0001	0.66
SOD1.1	SOD1	0.0245	0.83	FDPS.1	FDPS	<0.0001	0.47
SNRK.1	SNRK	<0.0001	0.35	FBXW7.1	FBXW7	<0.0001	0.66
SNAI1.1	SNAI1	<0.0001	0.70	fas.1	FAS	<0.0001	0.66
MADH4.1	SMAD4	<0.0001	0.47	ESRRG.3	ESRRG	0.0001	0.72
MADH2.1	SMAD2	<0.0001	0.50	ERG.1	ERG	<0.0001	0.44
SLC9A1.1	SLC9A1	0.0207	0.82	ERCC1.2	ERCC1	<0.0001	0.60
SLC34A1.1	SLC34A1	<0.0001	0.63	ERBB4.3	ERBB4	0.0018	0.74
SLC22A6.1	SLC22A6	0.0010	0.76	ErbB3.1	ERBB3	0.0031	0.79
SKIL.1	SKIL	<0.0001	0.64	HER2.3	ERBB2	<0.0001	0.53
PTPNS1.1	SIRPA	0.0075	0.81	EPHB4.1	EPHB4	<0.0001	0.51
SHANK3.1	SHANK3	<0.0001	0.37	EPHA2.1	EPHA2	<0.0001	0.40
SGK.1	SGK1	<0.0001	0.55	EPAS1.1	EPAS1	<0.0001	0.38
FRP1.3	SFRP1	0.0156	0.82	ENPP2.1	ENPP2	0.0001	0.72
SEMA3F.3	SEMA3F	<0.0001	0.50	ENPEP.1	ENPEP	<0.0001	0.65
SELPLG.1	SELPLG	0.0022	0.78	CD105.1	ENG	<0.0001	0.38
SELENBP1.1	SELENBP1	0.0003	0.75	EMP1.1	EMP1	<0.0001	0.64

Table 4b: Proxy genes for which increased expression is associated with lower tumor stage (p-value $\leq .05$)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
SDPR.1	SDPR	<0.0001	0.42	EMCN.1	EMCN	<0.0001	0.27
SDHA.1	SDHA	<0.0001	0.65	ELTD1.1	ELTD1	<0.0001	0.67
SCNN1A.2	SCNN1A	0.0024	0.77	EIF2C1.1	EIF2C1	<0.0001	0.51
SCN4B.1	SCN4B	<0.0001	0.50	EGR1.1	EGR1	<0.0001	0.59
S100A2.1	S100A2	0.0008	0.75	EGLN3.1	EGLN3	0.0002	0.75
KIAA1303 raptor.1	RPTOR	<0.0001	0.60	EGFR.2	EGFR	0.0072	0.81
RPS6KB1.3	RPS6KB1	<0.0001	0.60	EGF.3	EGF	0.0051	0.77
RPS6KA1.1	RPS6KA1	0.0002	0.71	EFNB2.1	EFNB2	<0.0001	0.45
RPS23.1	RPS23	0.0002	0.73	EFNB1.2	EFNB1	<0.0001	0.55
ROCK2.1	ROCK2	<0.0001	0.52	EEF1A1.1	EEF1A1	<0.0001	0.55
ROCK1.1	ROCK1	<0.0001	0.37	EDNRB.1	EDNRB	<0.0001	0.44
RIPK1.1	RIPK1	<0.0001	0.55	EDN2.1	EDN2	0.0012	0.74
rhoC.1	RHOC	<0.0001	0.66	EDN1 endothelin.1	EDN1	<0.0001	0.53
RhoB.1	RHOB	<0.0001	0.57	EBAG9.1	EBAG9	0.0240	0.83
ARHA.1	RHOA	<0.0001	0.50	DUSP1.1	DUSP1	0.0130	0.82
RHEB.2	RHEB	<0.0001	0.61	DPYS.1	DPYS	0.0355	0.85
RGS5.1	RGS5	<0.0001	0.37	DPEP1.1	DPEP1	<0.0001	0.66
FLJ22655.1	RERGL	<0.0001	0.33	DLL4.1	DLL4	<0.0001	0.66
RB1.1	RB1	<0.0001	0.61	DLC1.1	DLC1	<0.0001	0.42
RASSF1.1	RASSF1	0.0002	0.75	DKFZP564O08 23.1	DKFZP564O 0823	<0.0001	0.51
RARB.2	RARB	<0.0001	0.37	DICER1.2	DICER1	<0.0001	0.50
RALBP1.1	RALBP1	<0.0001	0.43	DIAPH1.1	DIAPH1	0.0219	0.83
RAF1.3	RAF1	<0.0001	0.59	DIABLO.1	DIABLO	0.0022	0.78
RAC1.3	RAC1	0.0356	0.84	DHPS.3	DHPS	<0.0001	0.55
PTPRG.1	PTPRG	<0.0001	0.35	DET1.1	DET1	0.0005	0.74
PTPRB.1	PTPRB	<0.0001	0.31	DEFB1.1	DEFB1	0.0002	0.73
PTN.1	PTN	<0.0001	0.56	DDC.1	DDC	<0.0001	0.72
PTK2.1	PTK2	<0.0001	0.45	DAPK1.3	DAPK1	<0.0001	0.42
PTHR1.1	PTH1R	<0.0001	0.45	CYR61.1	CYR61	<0.0001	0.59
PTEN.2	PTEN	<0.0001	0.51	CXCL12.1	CXCL12	<0.0001	0.62
PSMB9.1	PSMB9	0.0139	0.82	CX3CR1.1	CX3CR1	<0.0001	0.46
PSMB8.1	PSMB8	0.0243	0.83	CX3CL1.1	CX3CL1	<0.0001	0.47
PSMA7.1	PSMA7	0.0101	0.81	CUL1.1	CUL1	<0.0001	0.64
PRSS8.1	PRSS8	0.0025	0.78	CUBN.1	CUBN	<0.0001	0.55

Table 4b: Proxy genes for which increased expression is associated with lower tumor stage (p-value $\leq .05$)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
PRPS2.1	PRPS2	0.0190	0.82	CTSS.1	CTSS	0.0007	0.76
PRKCH.1	PRKCH	<0.0001	0.48	CTSH.2	CTSH	<0.0001	0.64
PRKCD.2	PRKCD	0.0001	0.72	B-Catenin.3	CTNNB1	<0.0001	0.54
PPP2CA.1	PPP2CA	<0.0001	0.58	A-Catenin.2	CTNNA1	<0.0001	0.65
PPARG.3	PPARG	<0.0001	0.40	CTGF.1	CTGF	<0.0001	0.71
PPAP2B.1	PPAP2B	<0.0001	0.31	CSF2RA.2	CSF2RA	0.0037	0.78
PLG.1	PLG	<0.0001	0.53	CSF1R.2	CSF1R	<0.0001	0.67
PLAT.1	PLAT	<0.0001	0.54	CSF1.1	CSF1	<0.0001	0.65
PLA2G4C.1	PLA2G4C	<0.0001	0.65	CRADD.1	CRADD	0.0032	0.79
PIK3CA.1	PIK3CA	<0.0001	0.53	COL4A2.1	COL4A2	<0.0001	0.65
PI3K.2	PIK3C2B	<0.0001	0.40	COL4A1.1	COL4A1	0.0067	0.80
PFKP.1	PFKP	0.0124	0.82	COL18A1.1	COL18A1	0.0001	0.72
CD31.3	PECAM1	<0.0001	0.42	CLU.3	CLU	0.0004	0.75
PDZK3.1	PDZK3	0.0003	0.72	CLDN10.1	CLDN10	<0.0001	0.65
PDZK1.1	PDZK1	<0.0001	0.58	CLCNKB.1	CLCNKB	<0.0001	0.52
PDGFRb.3	PDGFRB	<0.0001	0.62	CFLAR.1	CFLAR	<0.0001	0.60
PDGFD.2	PDGFD	<0.0001	0.43	CEACAM1.1	CEACAM1	<0.0001	0.55
PDGFC.3	PDGFC	<0.0001	0.51	p21.3	CDKN1A	0.0002	0.73
PDGFB.3	PDGFB	<0.0001	0.40	CDH6.1	CDH6	<0.0001	0.72
PDGFA.3	PDGFA	<0.0001	0.57	CDH5.1	CDH5	<0.0001	0.44
PCK1.1	PCK1	<0.0001	0.60	CDH2.1	CDH2	0.0392	0.85
PCCA.1	PCCA	<0.0001	0.58	CDH16.1	CDH16	<0.0001	0.70
PARD6A.1	PARD6A	0.0042	0.79	CDH13.1	CDH13	<0.0001	0.53
Pak1.2	PAK1	<0.0001	0.69	CDC25B.1	CDC25B	0.0037	0.79
PAH.1	PAH	0.0309	0.84	CD4.1	CD4	<0.0001	0.72
OGG1.1	OGG1	0.0024	0.78	CD36.1	CD36	<0.0001	0.51
BFGF.3	NUDT6	<0.0001	0.46	CD34.1	CD34	<0.0001	0.48
NPR1.1	NPR1	<0.0001	0.58	CD24.1	CD24	0.0206	0.83
NPM1.2	NPM1	<0.0001	0.65	CD14.1	CD14	0.0152	0.82
NOTCH3.1	NOTCH3	<0.0001	0.58	CCND1.3	CCND1	<0.0001	0.51
NOTCH2.1	NOTCH2	<0.0001	0.64	CCL4.2	CCL4	0.0017	0.77
NOTCH1.1	NOTCH1	<0.0001	0.44	MCP1.1	CCL2	<0.0001	0.66
NOS3.1	NOS3	<0.0001	0.44	CAT.1	CAT	<0.0001	0.53
NOS2A.3	NOS2	<0.0001	0.49	CASP10.1	CASP10	0.0001	0.72
NOL3.1	NOL3	0.0003	0.76	CALD1.2	CALD1	<0.0001	0.55

Table 4b: Proxy genes for which increased expression is associated with lower tumor stage (p-value $\leq .05$)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
NFX1.1	NFX1	<0.0001	0.50	CACNA2D1.1	CACNA2D1	0.0352	0.84
NFKBp50.3	NFKB1	<0.0001	0.59	CA9.3	CA9	0.0299	0.84
NFATC2.1	NFATC2	<0.0001	0.64	CA2.1	CA2	<0.0001	0.58
NFAT5.1	NFAT5	<0.0001	0.65	C3AR1.1	C3AR1	0.0010	0.77
MYRIP.2	MYRIP	0.0004	0.72	ECRG4.1	C2orf40	<0.0001	0.47
MYH11.1	MYH11	<0.0001	0.50	C1QA.1	C1QA	0.0119	0.82
cMYC.3	MYC	<0.0001	0.70	C13orf15.1	C13orf15	<0.0001	0.37
MX1.1	MX1	0.0103	0.81	BUB3.1	BUB3	<0.0001	0.67
MVP.1	MVP	0.0002	0.74	BTRC.1	BTRC	<0.0001	0.65
MUC1.2	MUC1	0.0005	0.75	CIAP1.2	BIRC2	<0.0001	0.64
FRAP1.1	MTOR	<0.0001	0.61	BIN1.3	BIN1	0.0001	0.73
MSH3.2	MSH3	<0.0001	0.60	BGN.1	BGN	<0.0001	0.63
MSH2.3	MSH2	<0.0001	0.53	Bclx.2	BCL2L1	<0.0001	0.58
STMY3.3	MMP11	0.0034	0.79	Bcl2.2	BCL2	<0.0001	0.37
GBL.1	MLST8	0.0011	0.77	Bax.1	BAX	0.0035	0.78
MIF.2	MIF	0.0008	0.76	Bak.2	BAK1	0.0215	0.83
MICA.1	MICA	<0.0001	0.70	BAG1.2	BAG1	<0.0001	0.40
MGMT.1	MGMT	<0.0001	0.60	ATP6V1B1.1	ATP6V1B1	<0.0001	0.54
MCM3.3	MCM3	<0.0001	0.68	ATP1A1.1	ATP1A1	0.0037	0.78
MCAM.1	MCAM	<0.0001	0.52	ASS1.1	ASS1	<0.0001	0.60
MARCKS.1	MARCKS	0.0001	0.73	ARRB1.1	ARRB1	<0.0001	0.45
ERK1.3	MAPK3	<0.0001	0.48	ARHGDIB.1	ARHGDIB	<0.0001	0.50
ERK2.3	MAPK1	0.0221	0.83	AQP1.1	AQP1	<0.0001	0.40
MAP4.1	MAP4	<0.0001	0.63	APOLD1.1	APOLD1	<0.0001	0.54
MAP2K3.1	MAP2K3	<0.0001	0.59	APC.4	APC	<0.0001	0.57
MAP2K1.1	MAP2K1	0.0002	0.74	ANXA4.1	ANXA4	0.0003	0.74
MAL2.1	MAL2	<0.0001	0.64	ANXA1.2	ANXA1	0.0001	0.73
MAL.1	MAL	<0.0001	0.49	ANTXR1.1	ANTXR1	0.0051	0.80
LYZ.1	LYZ	0.0318	0.84	ANGPTL4.1	ANGPTL4	0.0041	0.80
LTF.1	LTF	0.0131	0.80	ANGPTL3.3	ANGPTL3	0.0258	0.82
LRP2.1	LRP2	<0.0001	0.63	ANGPTL2.1	ANGPTL2	0.0015	0.77
LMO2.1	LMO2	<0.0001	0.56	ANGPT2.1	ANGPT2	<0.0001	0.72
LDB2.1	LDB2	<0.0001	0.41	ANGPT1.1	ANGPT1	<0.0001	0.45
LDB1.2	LDB1	<0.0001	0.54	ALDOB.1	ALDOB	<0.0001	0.60
LAMA4.1	LAMA4	0.0004	0.75	ALDH6A1.1	ALDH6A1	<0.0001	0.55

Table 4b: Proxy genes for which increased expression is associated with lower tumor stage (p-value $\leq .05$)

Gene	Official Symbol	Stage 3 vs. 1		Gene	Official Symbol	Stage 3 vs. 1	
		p-value	OR			p-value	OR
KRT7.1	KRT7	<0.0001	0.60	ALDH4.2	ALDH4A1	0.0124	0.82
K-ras.10	KRAS	<0.0001	0.68	AKT3.2	AKT3	<0.0001	0.43
KL.1	KL	<0.0001	0.49	AKT2.3	AKT2	<0.0001	0.64
Kitlng.4	KITLG	<0.0001	0.43	AKT1.3	AKT1	<0.0001	0.58
c-kit.2	KIT	<0.0001	0.60	AIF1.1	AIF1	0.0002	0.74
KDR.6	KDR	<0.0001	0.36	AHR.1	AHR	<0.0001	0.59
KCNJ15.1	KCNJ15	<0.0001	0.54	AGTR1.1	AGTR1	<0.0001	0.36
HTATIP.1	KAT5	<0.0001	0.40	ADH1B.1	ADH1B	0.0015	0.77
G-Catenin.1	JUP	<0.0001	0.42	ADD1.1	ADD1	<0.0001	0.40
AP-1 (JUN official).2	JUN	0.0001	0.73	ADAMTS5.1	ADAMTS5	0.0006	0.73
JAG1.1	JAG1	<0.0001	0.42	ADAM17.1	ADAM17	<0.0001	0.72
ITGB5.1	ITGB5	0.0115	0.81	ACE2.1	ACE2	<0.0001	0.61
ITGB1.1	ITGB1	<0.0001	0.64	ACADSB.1	ACADSB	<0.0001	0.54
ITGA7.1	ITGA7	<0.0001	0.54	BCRP.1	ABCG2	<0.0001	0.41
ITGA6.2	ITGA6	<0.0001	0.51	MRP4.2	ABCC4	<0.0001	0.65
ITGA5.1	ITGA5	0.0325	0.84	MRP3.1	ABCC3	0.0005	0.76
ITGA4.2	ITGA4	<0.0001	0.54	MRP1.1	ABCC1	0.0017	0.78
ITGA3.2	ITGA3	<0.0001	0.61	ABCB1.5	ABCB1	0.0003	0.75
IQGAP2.1	IQGAP2	<0.0001	0.63	NPD009 (ABAT official).3	ABAT	<0.0001	0.70
INSR.1	INSR	<0.0001	0.59	AAMP.1	AAMP	0.0292	0.84
IMP3.1	IMP3	<0.0001	0.54	A2M.1	A2M	<0.0001	0.36

Table 5a: Proxy genes for which increased expression is associated with higher tumor grade (p-value $\leq .05$)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
WT1.1	WT1	<0.0001	1.39	HPD.1	HPD	0.0054	1.20
VTN.1	VTN	0.0359	1.16	HIST1H1D.1	HIST1H1D	0.0248	1.16
VDR.2	VDR	0.0001	1.29	HGD.1	HGD	<0.0001	1.41
UBE2T.1	UBE2T	<0.0001	1.81	GZMA.1	GZMA	0.0006	1.26
TP.3	TYMP	<0.0001	1.35	GPX2.2	GPX2	0.0182	1.18
C20 orf1.1	TPX2	<0.0001	2.14	GPX1.2	GPX1	0.0008	1.25
TOP2A.4	TOP2A	<0.0001	2.07	FCGR3A.1	FCGR3A	0.0003	1.27
TNFSF13B.1	TNFSF13B	0.0062	1.20	fasl.2	FASLG	0.0045	1.21
TK1.2	TK1	<0.0001	1.66	FABP1.1	FABP1	<0.0001	1.32
TGFBI.1	TGFBI	0.0452	1.14	F2.1	F2	<0.0001	1.77
STAT1.3	STAT1	<0.0001	1.31	ESPL1.3	ESPL1	<0.0001	1.60
SQSTM1.1	SQSTM1	0.0003	1.27	E2F1.3	E2F1	<0.0001	1.36
OPN, osteopontin.3	SPP1	0.0002	1.29	CXCR6.1	CXCR6	<0.0001	1.50
SLC7A5.2	SLC7A5	0.0002	1.28	BLR1.1	CXCR5	0.0338	1.15
SLC16A3.1	SLC16A3	0.0052	1.21	CXCL9.1	CXCL9	0.0001	1.29
SLC13A3.1	SLC13A3	0.0003	1.27	CXCL10.1	CXCL10	0.0078	1.19
SFN.1	SFN	0.0066	1.20	GRO1.2	CXCL1	<0.0001	1.39
SEMA3C.1	SEMA3C	<0.0001	1.32	CTSD.2	CTSD	0.0183	1.17
SAA2.2	SAA2	<0.0001	2.13	CTSB.1	CTSB	0.0006	1.26
S100A1.1	S100A1	<0.0001	1.40	CRP.1	CRP	0.0342	1.15
RRM2.1	RRM2	<0.0001	1.71	CP.1	CP	<0.0001	1.37
RPLP1.1	RPLP1	0.0007	1.25	Chk2.3	CHEK2	<0.0001	1.37
RAD51.1	RAD51	<0.0001	1.53	Chk1.2	CHEK1	<0.0001	1.37
PTTG1.2	PTTG1	<0.0001	1.89	CENPF.1	CENPF	<0.0001	1.78
PSMB9.1	PSMB9	0.0010	1.25	CD8A.1	CD8A	<0.0001	1.35
PSMB8.1	PSMB8	0.0181	1.17	CD82.3	CD82	<0.0001	1.50
PRKCB1.1	PRKCB	0.0218	1.16	TNFSF7.1	CD70	<0.0001	1.43
PDCD1.1	PDCD1	<0.0001	1.43	CCNE1.1	CCNE1	0.0002	1.29
PCSK6.1	PCSK6	0.0009	1.25	CCNB1.2	CCNB1	<0.0001	1.75
PCNA.2	PCNA	0.0041	1.21	CCL5.2	CCL5	<0.0001	1.63
NME2.1	NME2	0.0106	1.19	CCL20.1	CCL20	0.0082	1.19
MYBL2.1	MYBL2	<0.0001	1.70	CAV2.1	CAV2	0.0210	1.17
MMP9.1	MMP9	<0.0001	1.46	CA12.1	CA12	<0.0001	1.41
MMP14.1	MMP14	0.0003	1.28	C3.1	C3	<0.0001	1.34
Ki-67.2	MKI67	<0.0001	1.70	C1QB.1	C1QB	0.0201	1.17

Table 5a: Proxy genes for which increased expression is associated with higher tumor grade (p-value $\leq$.05)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
mGST1.2	MGST1	<0.0001	2.13	BUB1.1	BUB1	<0.0001	2.16
cMet.2	MET	<0.0001	1.57	BRCA1.2	BRCA1	0.0004	1.26
MDK.1	MDK	<0.0001	1.55	SURV.2	BIRC5	<0.0001	1.93
MDH2.1	MDH2	<0.0001	1.35	clAP2.2	BIRC3	<0.0001	1.44
MCM2.2	MCM2	<0.0001	1.33	STK15.2	AURKA	<0.0001	1.38
LOX.1	LOX	<0.0001	1.35	ATP5E.1	ATP5E	<0.0001	1.45
LMNB1.1	LMNB1	<0.0001	1.76	APOL1.1	APOL1	<0.0001	1.47
LIMK1.1	LIMK1	<0.0001	1.43	APOE.1	APOE	<0.0001	1.40
LAPTM5.1	LAPTM5	0.0044	1.21	APOC1.3	APOC1	<0.0001	1.42
LAMB3.1	LAMB3	<0.0001	1.49	ANXA2.2	ANXA2	0.0020	1.23
L1CAM.1	L1CAM	0.0338	1.15	ANGPTL3.3	ANGPTL3	<0.0001	1.32
KLRK1.2	KLRK1	0.0412	1.15	AMACR1.1	AMACR	0.0382	1.15
CD18.2	ITGB2	0.0069	1.21	ALOX5.1	ALOX5	0.0001	1.29
IL-8.1	IL8	0.0088	1.19	ALDH4.2	ALDH4A1	0.0002	1.28
IL6.3	IL6	0.0091	1.19	ADAM8.1	ADAM8	0.0006	1.26
ICAM1.1	ICAM1	0.0014	1.24	MRP2.3	ABCC2	<0.0001	1.97
HSPA8.1	HSPA8	<0.0001	1.39				

Table 5b: Proxy genes for which increased expression is associated with lower tumor grade (p-value $\leq .05$)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
ZHX2.1	ZHX2	0.0061	0.83	ITGA4.2	ITGA4	<0.0001	0.75
YB-1.2	YBX1	<0.0001	0.62	ITGA3.2	ITGA3	0.0128	0.85
XPNPEP2.2	XPNPEP2	0.0040	0.82	IQGAP2.1	IQGAP2	<0.0001	0.68
XIAP.1	XIAP	<0.0001	0.64	INSR.1	INSR	<0.0001	0.46
WISP1.1	WISP1	<0.0001	0.58	INHBA.1	INHBA	0.0049	0.83
VWF.1	VWF	<0.0001	0.34	IMP3.1	IMP3	<0.0001	0.67
VHL.1	VHL	0.0088	0.84	IL6ST.3	IL6ST	<0.0001	0.43
VEGF.1	VEGFA	<0.0001	0.48	IL1B.1	IL1B	0.0102	0.84
VCAN.1	VCAN	0.0023	0.82	IL15.1	IL15	0.0001	0.76
VCAM1.1	VCAM1	0.0049	0.83	IL10.3	IL10	0.0239	0.86
USP34.1	USP34	<0.0001	0.62	IGFBP6.1	IGFBP6	<0.0001	0.75
UMOD.1	UMOD	<0.0001	0.69	IGFBP5.1	IGFBP5	<0.0001	0.74
UGCG.1	UGCG	<0.0001	0.58	IGFBP2.1	IGFBP2	<0.0001	0.76
UBB.1	UBB	<0.0001	0.62	IGF2.2	IGF2	<0.0001	0.69
UBE1C.1	UBA3	<0.0001	0.67	IGF1R.3	IGF1R	<0.0001	0.43
tusc4.2	TUSC4	<0.0001	0.61	ID3.1	ID3	<0.0001	0.48
TUSC2.1	TUSC2	0.0481	0.88	ID2.4	ID2	<0.0001	0.64
TSPAN7.2	TSPAN7	<0.0001	0.29	ID1.1	ID1	<0.0001	0.37
TSC2.1	TSC2	<0.0001	0.60	ICAM2.1	ICAM2	<0.0001	0.42
TSC1.1	TSC1	<0.0001	0.52	HYAL2.1	HYAL2	<0.0001	0.32
P53.2	TP53	<0.0001	0.66	HYAL1.1	HYAL1	<0.0001	0.52
TOP2B.2	TOP2B	<0.0001	0.68	HSPG2.1	HSPG2	<0.0001	0.33
TNIP2.1	TNIP2	0.0001	0.76	HSPA1A.1	HSPA1A	0.0022	0.81
TNFSF12.1	TNFSF12	<0.0001	0.54	HSP90AB1.1	HSP90AB1	<0.0001	0.68
TNFRSF11B.1	TNFRSF11B	<0.0001	0.73	HSD11B2.1	HSD11B2	<0.0001	0.43
TNFRSF10D.1	TNFRSF10D	<0.0001	0.51	HPCAL1.1	HPCAL1	<0.0001	0.69
TNFRSF10C.3	TNFRSF10C	0.0003	0.78	HNRPAB.3	HNRNPAB	0.0432	0.87
DR5.2	TNFRSF10B	<0.0001	0.69	HMGB1.1	HMGB1	<0.0001	0.47
TNFAIP6.1	TNFAIP6	0.0338	0.87	HIF1AN.1	HIF1AN	<0.0001	0.64
TNFAIP3.1	TNFAIP3	0.0083	0.84	HIF1A.3	HIF1A	<0.0001	0.55
TNF.1	TNF	0.0392	0.87	HGF.4	HGF	0.0022	0.81
TMEM47.1	TMEM47	<0.0001	0.33	HDAC1.1	HDAC1	<0.0001	0.48
TMEM27.1	TMEM27	<0.0001	0.73	HADH.1	HADH	<0.0001	0.63
TIMP3.3	TIMP3	<0.0001	0.29	GSTp.3	GSTP1	<0.0001	0.66

Table 5b: Proxy genes for which increased expression is associated with lower tumor grade (p-value $\leq .05$)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
TIMP2.1	TIMP2	<0.0001	0.54	GSTM3.2	GSTM3	<0.0001	0.53
THBS1.1	THBS1	<0.0001	0.58	GSTM1.1	GSTM1	<0.0001	0.61
THBD.1	THBD	<0.0001	0.66	GRB7.2	GRB7	<0.0001	0.73
TGFBR2.3	TGFBR2	<0.0001	0.32	GRB14.1	GRB14	0.0001	0.76
TGFBR1.1	TGFBR1	<0.0001	0.59	GPC3.1	GPC3	0.0012	0.80
TGFB2.2	TGFB2	<0.0001	0.54	GNAS.1	GNAS	<0.0001	0.70
TGFb1.1	TGFB1	<0.0001	0.66	GMNN.1	GMNN	0.0006	0.80
TGFA.2	TGFA	<0.0001	0.68	GJA1.1	GJA1	<0.0001	0.66
TEK.1	TEK	<0.0001	0.34	GCLM.2	GCLM	0.0319	0.87
cripto (TDGF1 official).1	TDGF1	0.0328	0.86	GCLC.3	GCLC	<0.0001	0.57
TCF4.1	TCF4	<0.0001	0.33	GATM.1	GATM	0.0006	0.79
TAGLN.1	TAGLN	<0.0001	0.58	GATA3.3	GATA3	0.0029	0.82
TACSTD2.1	TACSTD2	<0.0001	0.61	GAS2.1	GAS2	0.0136	0.84
SUCLG1.1	SUCLG1	<0.0001	0.76	GADD45B.1	GADD45B	<0.0001	0.54
STK11.1	STK11	<0.0001	0.53	FST.1	FST	0.0197	0.85
STC2.1	STC2	0.0085	0.84	FOS.1	FOS	<0.0001	0.44
STAT5B.2	STAT5B	<0.0001	0.40	FOLR1.1	FOLR1	<0.0001	0.75
STAT5A.1	STAT5A	<0.0001	0.60	FLT4.1	FLT4	<0.0001	0.34
STAT3.1	STAT3	<0.0001	0.48	FLT3LG.1	FLT3LG	<0.0001	0.73
SPRY1.1	SPRY1	<0.0001	0.40	FLT1.1	FLT1	<0.0001	0.29
SPAST.1	SPAST	0.0002	0.78	FILIP1.1	FILIP1	<0.0001	0.47
SPARCL1.1	SPARCL1	<0.0001	0.41	FIGF.1	FIGF	<0.0001	0.70
SPARC.1	SPARC	<0.0001	0.50	FHL1.1	FHL1	<0.0001	0.54
SNRK.1	SNRK	<0.0001	0.29	FHIT.1	FHIT	<0.0001	0.72
SNAI1.1	SNAI1	<0.0001	0.54	FH.1	FH	0.0203	0.86
MADH4.1	SMAD4	<0.0001	0.39	FGFR2 isoform 1.1	FGFR2	<0.0001	0.62
MADH2.1	SMAD2	<0.0001	0.40	FGFR1.3	FGFR1	<0.0001	0.48
SLC9A1.1	SLC9A1	<0.0001	0.75	FGF2.2	FGF2	<0.0001	0.62
SLC34A1.1	SLC34A1	0.0001	0.76	FGF1.1	FGF1	<0.0001	0.61
SKIL.1	SKIL	<0.0001	0.52	FDPS.1	FDPS	<0.0001	0.52
SHC1.1	SHC1	0.0063	0.83	FBXW7.1	FBXW7	<0.0001	0.57
SHANK3.1	SHANK3	<0.0001	0.27	FAP.1	FAP	0.0440	0.87
SGK.1	SGK1	<0.0001	0.60	ESRRG.3	ESRRG	0.0340	0.87
FRP1.3	SFRP1	0.0003	0.78	ERG.1	ERG	<0.0001	0.36

Table 5b: Proxy genes for which increased expression is associated with lower tumor grade (p-value $\leq$.05)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
PAI1.3	SERPINE1	0.0115	0.85	ERCC4.1	ERCC4	0.0337	0.87
SEMA3F.3	SEMA3F	<0.0001	0.45	ERCC1.2	ERCC1	<0.0001	0.59
SELENBP1.1	SELENBP1	<0.0001	0.63	ERBB4.3	ERBB4	<0.0001	0.66
SELE.1	SELE	<0.0001	0.74	HER2.3	ERBB2	<0.0001	0.57
SDPR.1	SDPR	<0.0001	0.35	EPHB4.1	EPHB4	<0.0001	0.43
SDHA.1	SDHA	<0.0001	0.69	EPHA2.1	EPHA2	<0.0001	0.44
SCNN1A.2	SCNN1A	0.0011	0.80	EPAS1.1	EPAS1	<0.0001	0.26
SCN4B.1	SCN4B	<0.0001	0.47	ENPP2.1	ENPP2	<0.0001	0.62
S100A2.1	S100A2	<0.0001	0.72	ENPEP.1	ENPEP	<0.0001	0.75
RUNX1.1	RUNX1	0.0001	0.77	ENO2.1	ENO2	0.0449	0.88
RRM1.2	RRM1	0.0438	0.87	CD105.1	ENG	<0.0001	0.30
KIAA1303 raptor.1	RPTOR	<0.0001	0.53	EMP1.1	EMP1	<0.0001	0.42
RPS6KB1.3	RPS6KB1	<0.0001	0.49	EMCN.1	EMCN	<0.0001	0.31
RPS23.1	RPS23	<0.0001	0.59	ELTD1.1	ELTD1	<0.0001	0.59
ROCK2.1	ROCK2	<0.0001	0.42	EIF2C1.1	EIF2C1	<0.0001	0.63
ROCK1.1	ROCK1	<0.0001	0.31	EGR1.1	EGR1	<0.0001	0.50
RIPK1.1	RIPK1	<0.0001	0.50	EGLN3.1	EGLN3	<0.0001	0.69
rhoC.1	RHOC	<0.0001	0.70	EGF.3	EGF	0.0200	0.85
RhoB.1	RHOB	<0.0001	0.36	EFNB2.1	EFNB2	<0.0001	0.36
ARHA.1	RHOA	<0.0001	0.34	EFNB1.2	EFNB1	<0.0001	0.46
RHEB.2	RHEB	<0.0001	0.72	EEF1A1.1	EEF1A1	<0.0001	0.39
RGS5.1	RGS5	<0.0001	0.30	EDNRB.1	EDNRB	<0.0001	0.33
FLJ22655.1	RERGL	<0.0001	0.42	EDN2.1	EDN2	<0.0001	0.67
NFKBp65.3	RELA	<0.0001	0.69	EDN1 endothelin.1	EDN1	<0.0001	0.41
RB1.1	RB1	<0.0001	0.74	EBAG9.1	EBAG9	0.0057	0.83
RASSF1.1	RASSF1	<0.0001	0.60	DUSP1.1	DUSP1	<0.0001	0.54
RARB.2	RARB	<0.0001	0.41	DPEP1.1	DPEP1	0.0001	0.76
RALBP1.1	RALBP1	<0.0001	0.47	DLL4.1	DLL4	<0.0001	0.49
RAF1.3	RAF1	<0.0001	0.48	DLC1.1	DLC1	<0.0001	0.33
RAC1.3	RAC1	<0.0001	0.75	DKFZP564O08 23.1	DKFZP564O 0823	<0.0001	0.46
PXDN.1	PXDN	<0.0001	0.74	DICER1.2	DICER1	<0.0001	0.41
PTPRG.1	PTPRG	<0.0001	0.29	DIAPH1.1	DIAPH1	0.0022	0.81
PTPRB.1	PTPRB	<0.0001	0.27	DIABLO.1	DIABLO	<0.0001	0.73
PTN.1	PTN	<0.0001	0.58	DHPS.3	DHPS	<0.0001	0.41

Table 5b: Proxy genes for which increased expression is associated with lower tumor grade (p-value $\leq .05$)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
PTK2.1	PTK2	<0.0001	0.36	DET1.1	DET1	<0.0001	0.63
PTHR1.1	PTH1R	<0.0001	0.50	DEFB1.1	DEFB1	0.0001	0.77
PTEN.2	PTEN	<0.0001	0.40	DAPK1.3	DAPK1	<0.0001	0.48
PSMA7.1	PSMA7	0.0002	0.78	DAG1.1	DAG1	0.0150	0.85
PRPS2.1	PRPS2	0.0012	0.80	CYR61.1	CYR61	<0.0001	0.49
PROM2.1	PROM2	0.0326	0.87	CXCL12.1	CXCL12	<0.0001	0.64
PRKCH.1	PRKCH	<0.0001	0.45	CX3CR1.1	CX3CR1	0.0008	0.80
PRKCD.2	PRKCD	<0.0001	0.76	CX3CL1.1	CX3CL1	<0.0001	0.55
PPP2CA.1	PPP2CA	<0.0001	0.68	CUL1.1	CUL1	<0.0001	0.62
PPARG.3	PPARG	<0.0001	0.42	CUBN.1	CUBN	0.0081	0.84
PPAP2B.1	PPAP2B	<0.0001	0.32	CTSL.2	CTSL1	0.0328	0.87
PMP22.1	PMP22	<0.0001	0.61	B-Catenin.3	CTNNB1	<0.0001	0.36
PLG.1	PLG	<0.0001	0.75	A-Catenin.2	CTNNA1	<0.0001	0.73
PLAT.1	PLAT	<0.0001	0.42	CTGF.1	CTGF	<0.0001	0.61
PLA2G4C.1	PLA2G4C	0.0002	0.78	CSF1.1	CSF1	<0.0001	0.66
PIK3CA.1	PIK3CA	<0.0001	0.48	CRADD.1	CRADD	0.0151	0.85
PI3K.2	PIK3C2B	<0.0001	0.53	COL5A2.2	COL5A2	0.0155	0.85
PGF.1	PGF	<0.0001	0.74	COL4A2.1	COL4A2	<0.0001	0.51
PFKP.1	PFKP	0.0008	0.80	COL4A1.1	COL4A1	<0.0001	0.59
CD31.3	PECAM1	<0.0001	0.32	COL1A2.1	COL1A2	0.0005	0.79
PDZK3.1	PDZK3	<0.0001	0.72	COL18A1.1	COL18A1	<0.0001	0.53
PDZK1.1	PDZK1	<0.0001	0.76	CLDN10.1	CLDN10	0.0351	0.87
PDGFRb.3	PDGFRB	<0.0001	0.42	CLCNKB.1	CLCNKB	<0.0001	0.73
PDGFRa.2	PDGFRA	0.0022	0.82	CFLAR.1	CFLAR	<0.0001	0.45
PDGFD.2	PDGFD	<0.0001	0.39	CEACAM1.1	CEACAM1	<0.0001	0.64
PDGFC.3	PDGFC	<0.0001	0.57	p27.3	CDKN1B	<0.0001	0.65
PDGFB.3	PDGFB	<0.0001	0.33	p21.3	CDKN1A	<0.0001	0.52
PDGFA.3	PDGFA	<0.0001	0.42	CDK4.1	CDK4	0.0200	0.86
PCK1.1	PCK1	<0.0001	0.71	CDH5.1	CDH5	<0.0001	0.31
PCCA.1	PCCA	<0.0001	0.75	CDH16.1	CDH16	0.0005	0.79
PARD6A.1	PARD6A	0.0088	0.84	CDH13.1	CDH13	<0.0001	0.40
Pak1.2	PAK1	0.0014	0.81	CD99.1	CD99	0.0001	0.77
PAH.1	PAH	0.0160	0.85	CD44.1	CD44_1	0.0216	0.86
OGG1.1	OGG1	0.0139	0.85	CD36.1	CD36	<0.0001	0.42
BFGF.3	NUDT6	<0.0001	0.63	CD34.1	CD34	<0.0001	0.39

Table 5b: Proxy genes for which increased expression is associated with lower tumor grade (p-value $\leq .05$)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
NPR1.1	NPR1	<0.0001	0.42	CD14.1	CD14	0.0021	0.81
NPM1.2	NPM1	<0.0001	0.61	CCND1.3	CCND1	<0.0001	0.63
NOTCH3.1	NOTCH3	<0.0001	0.40	MCP1.1	CCL2	<0.0001	0.69
NOTCH2.1	NOTCH2	<0.0001	0.58	CAT.1	CAT	0.0014	0.81
NOTCH1.1	NOTCH1	<0.0001	0.38	CASP10.1	CASP10	<0.0001	0.65
NOS3.1	NOS3	<0.0001	0.42	CALD1.2	CALD1	<0.0001	0.44
NOS2A.3	NOS2	<0.0001	0.56	CACNA2D1.1	CACNA2D1	0.0003	0.78
NOL3.1	NOL3	<0.0001	0.61	CA9.3	CA9	0.0269	0.86
NFX1.1	NFX1	<0.0001	0.53	CA2.1	CA2	0.0001	0.77
NFKBp50.3	NFKB1	<0.0001	0.59	C7.1	C7	<0.0001	0.71
NFATC2.1	NFATC2	<0.0001	0.73	C3AR1.1	C3AR1	0.0032	0.82
NFAT5.1	NFAT5	<0.0001	0.58	ECRG4.1	C2orf40	<0.0001	0.50
MYRIP.2	MYRIP	<0.0001	0.63	C13orf15.1	C13orf15	<0.0001	0.37
MYH11.1	MYH11	<0.0001	0.48	BUB3.1	BUB3	<0.0001	0.65
cMYC.3	MYC	<0.0001	0.68	BTRC.1	BTRC	<0.0001	0.61
MX1.1	MX1	0.0087	0.84	BNIP3.1	BNIP3	0.0018	0.81
MVP.1	MVP	0.0291	0.87	CIAP1.2	BIRC2	<0.0001	0.61
MUC1.2	MUC1	0.0043	0.83	BGN.1	BGN	<0.0001	0.43
FRAP1.1	MTOR	<0.0001	0.61	Bclx.2	BCL2L1	<0.0001	0.76
MT1X.1	MT1X	0.0276	0.86	Bcl2.2	BCL2	<0.0001	0.45
MSH3.2	MSH3	0.0001	0.76	BAG1.2	BAG1	<0.0001	0.47
MSH2.3	MSH2	<0.0001	0.60	AXL.1	AXL	<0.0001	0.74
MMP2.2	MMP2	<0.0001	0.74	ATP6V1B1.1	ATP6V1B1	<0.0001	0.65
STMY3.3	MMP11	<0.0001	0.70	ASS1.1	ASS1	<0.0001	0.67
MIF.2	MIF	0.0004	0.79	ARRB1.1	ARRB1	<0.0001	0.49
MICA.1	MICA	<0.0001	0.60	ARHGDIB.1	ARHGDIB	<0.0001	0.50
MGMT.1	MGMT	<0.0001	0.57	ARF1.1	ARF1	<0.0001	0.69
MCM3.3	MCM3	<0.0001	0.73	AREG.2	AREG	0.0007	0.80
MCAM.1	MCAM	<0.0001	0.42	AQP1.1	AQP1	<0.0001	0.49
MARCKS.1	MARCKS	<0.0001	0.63	APOLD1.1	APOLD1	<0.0001	0.40
ERK1.3	MAPK3	<0.0001	0.35	APC.4	APC	<0.0001	0.62
ERK2.3	MAPK1	<0.0001	0.71	APAF1.2	APAF1	<0.0001	0.76
MAP4.1	MAP4	<0.0001	0.58	ANXA1.2	ANXA1	<0.0001	0.65
MAP2K3.1	MAP2K3	<0.0001	0.60	ANTXR1.1	ANTXR1	<0.0001	0.61
MAP2K1.1	MAP2K1	<0.0001	0.62	ANGPTL4.1	ANGPTL4	<0.0001	0.75

Table 5b: Proxy genes for which increased expression is associated with lower tumor grade (p-value $\leq$.05)

Gene	Official Symbol	CCF Grade		Gene	Official Symbol	CCF Grade	
		p-value	OR			p-value	OR
MAL.1	MAL	<0.0001	0.63	ANGPTL2.1	ANGPTL2	<0.0001	0.60
LRP2.1	LRP2	0.0275	0.86	ANGPT2.1	ANGPT2	<0.0001	0.61
LMO2.1	LMO2	<0.0001	0.68	ANGPT1.1	ANGPT1	<0.0001	0.33
LDB2.1	LDB2	<0.0001	0.28	ALDOB.1	ALDOB	0.0348	0.87
LDB1.2	LDB1	<0.0001	0.52	ALDH6A1.1	ALDH6A1	<0.0001	0.61
LAMB1.1	LAMB1	<0.0001	0.73	AKT3.2	AKT3	<0.0001	0.30
LAMA4.1	LAMA4	<0.0001	0.55	AKT2.3	AKT2	<0.0001	0.65
KRT7.1	KRT7	<0.0001	0.61	AKT1.3	AKT1	<0.0001	0.50
K-ras.10	KRAS	<0.0001	0.54	AHR.1	AHR	<0.0001	0.55
KL.1	KL	<0.0001	0.62	AGTR1.1	AGTR1	<0.0001	0.44
Kitlng.4	KITLG	<0.0001	0.46	ADH1B.1	ADH1B	<0.0001	0.70
c-kit.2	KIT	<0.0001	0.48	ADD1.1	ADD1	<0.0001	0.40
KDR.6	KDR	<0.0001	0.33	ADAMTS9.1	ADAMTS9	<0.0001	0.64
KCNJ15.1	KCNJ15	<0.0001	0.67	ADAMTS5.1	ADAMTS5	<0.0001	0.56
HTATIP.1	KAT5	<0.0001	0.33	ADAMTS4.1	ADAMTS4	<0.0001	0.66
G-Catenin.1	JUP	<0.0001	0.39	ADAMTS2.1	ADAMTS2	<0.0001	0.69
AP-1 (JUN official).2	JUN	<0.0001	0.54	ADAMTS1.1	ADAMTS1	<0.0001	0.51
JAG1.1	JAG1	<0.0001	0.28	ADAM17.1	ADAM17	<0.0001	0.67
ITGB5.1	ITGB5	<0.0001	0.60	ACADSB.1	ACADSB	<0.0001	0.63
ITGB3.1	ITGB3	<0.0001	0.68	BCRP.1	ABCG2	<0.0001	0.43
ITGB1.1	ITGB1	<0.0001	0.46	MRP4.2	ABCC4	0.0337	0.87
ITGA7.1	ITGA7	<0.0001	0.40	AAMP.1	AAMP	<0.0001	0.71
ITGA6.2	ITGA6	<0.0001	0.51	A2M.1	A2M	<0.0001	0.29
ITGA5.1	ITGA5	<0.0001	0.62				

Table 6a: Proxy genes for which increased expression is associated with the presence of necrosis (p-value $\leq .05$)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
WT1.1	WT1	<0.0001	1.57	KRT19.3	KRT19	0.0001	1.43
VTN.1	VTN	<0.0001	1.38	ITGB4.2	ITGB4	0.0492	1.19
VDR.2	VDR	0.0013	1.34	ISG20.1	ISG20	0.0006	1.38
UBE2T.1	UBE2T	<0.0001	2.08	IL-8.1	IL8	<0.0001	2.40
TP.3	TYMP	0.0008	1.37	IL6.3	IL6	<0.0001	2.02
TSPAN8.1	TSPAN8	0.0016	1.29	ICAM1.1	ICAM1	<0.0001	1.75
C20 orf1.1	TPX2	<0.0001	2.64	HSPA8.1	HSPA8	0.0004	1.38
TOP2A.4	TOP2A	<0.0001	2.02	HIST1H1D.1	HIST1H1D	0.0017	1.33
TNFSF13B.1	TNFSF13B	0.0002	1.38	GPX1.2	GPX1	<0.0001	1.46
TK1.2	TK1	<0.0001	1.71	FZD2.2	FZD2	0.0031	1.27
TIMP1.1	TIMP1	0.0076	1.27	FN1.1	FN1	<0.0001	1.45
TGFBI.1	TGFBI	<0.0001	1.69	FCGR3A.1	FCGR3A	0.0001	1.43
OPN, osteopontin.3	SPP1	<0.0001	1.81	FCER1G.2	FCER1G	<0.0001	1.50
SPHK1.1	SPHK1	<0.0001	1.44	FAP.1	FAP	0.0395	1.20
SLC7A5.2	SLC7A5	<0.0001	2.12	F3.1	F3	<0.0001	1.62
SLC2A1.1	SLC2A1	<0.0001	1.46	F2.1	F2	<0.0001	1.76
SLC16A3.1	SLC16A3	0.0001	1.48	ESPL1.3	ESPL1	0.0008	1.33
SLC13A3.1	SLC13A3	0.0019	1.29	EPHB2.1	EPHB2	0.0037	1.28
SHC1.1	SHC1	0.0156	1.24	EPHB1.3	EPHB1	0.0041	1.25
SFN.1	SFN	<0.0001	1.57	EPB41L3.1	EPB41L3	0.0410	1.19
PAI1.3	SERPINE1	0.0164	1.25	ENO2.1	ENO2	0.0001	1.46
SERPINA5.1	SERPINA5	0.0016	1.27	EIF4EBP1.1	EIF4EBP1	<0.0001	1.60
SEMA3C.1	SEMA3C	<0.0001	2.14	E2F1.3	E2F1	<0.0001	1.65
SELL.1	SELL	0.0096	1.26	CXCR6.1	CXCR6	<0.0001	1.48
SAA2.2	SAA2	<0.0001	2.50	CXCR4.3	CXCR4	0.0118	1.26
RRM2.1	RRM2	<0.0001	1.86	GRO1.2	CXCL1	<0.0001	1.83
RPLP1.1	RPLP1	0.0025	1.32	CTSB.1	CTSB	<0.0001	1.70
RND3.1	RND3	0.0002	1.41	CRP.1	CRP	0.0494	1.16
RAD51.1	RAD51	<0.0001	1.51	CP.1	CP	<0.0001	1.98
PTTG1.2	PTTG1	<0.0001	2.52	COL7A1.1	COL7A1	<0.0001	1.47
COX2.1	PTGS2	0.0002	1.36	COL1A1.1	COL1A1	0.0059	1.28
PRKCB1.1	PRKCB	0.0188	1.23	Chk2.3	CHEK2	0.0010	1.33
PRKCA.1	PRKCA	0.0339	1.21	Chk1.2	CHEK1	<0.0001	1.49
PLAUR.3	PLAUR	<0.0001	1.60	CENPF.1	CENPF	<0.0001	2.08

Table 6a: Proxy genes for which increased expression is associated with the presence of necrosis (p-value $\leq$.05)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
upa.3	PLAU	0.0002	1.41	CD82.3	CD82	<0.0001	2.09
PF4.1	PF4	0.0003	1.34	CD68.2	CD68	0.0163	1.24
PDCD1.1	PDCD1	<0.0001	1.40	CD44s.1	CD44_s	<0.0001	1.66
PCSK6.1	PCSK6	0.0297	1.22	CCNE2.2	CCNE2_2	<0.0001	1.50
PCNA.2	PCNA	0.0025	1.33	CCNE1.1	CCNE1	<0.0001	1.44
NNMT.1	NNMT	0.0206	1.22	CCNB1.2	CCNB1	<0.0001	2.28
NME2.1	NME2	0.0124	1.25	CCL5.2	CCL5	<0.0001	1.45
MYBL2.1	MYBL2	<0.0001	1.90	CCL20.1	CCL20	0.0121	1.24
MT1X.1	MT1X	0.0003	1.39	CAV2.1	CAV2	0.0003	1.35
MMP9.1	MMP9	<0.0001	1.96	CA12.1	CA12	<0.0001	2.11
MMP7.1	MMP7	<0.0001	1.50	C3.1	C3	<0.0001	1.58
MMP14.1	MMP14	<0.0001	1.50	C1QB.1	C1QB	0.0032	1.31
Ki-67.2	MKI67	<0.0001	1.96	BUB1.1	BUB1	<0.0001	2.25
mGST1.2	MGST1	<0.0001	1.63	BRCA1.2	BRCA1	0.0006	1.35
cMet.2	MET	0.0357	1.22	SURV.2	BIRC5	<0.0001	2.11
MDK.1	MDK	<0.0001	1.78	cIAP2.2	BIRC3	<0.0001	1.47
MCM2.2	MCM2	0.0003	1.40	BCL2A1.1	BCL2A1	0.0004	1.34
LRRC2.1	LRRC2	0.0114	1.22	STK15.2	AURKA	<0.0001	1.61
LOX.1	LOX	<0.0001	1.99	PRO2000.3	ATAD2	0.0166	1.24
LMNB1.1	LMNB1	<0.0001	2.04	APOL1.1	APOL1	<0.0001	1.54
LIMK1.1	LIMK1	<0.0001	2.51	APOC1.3	APOC1	0.0026	1.30
LGALS9.1	LGALS9	0.0136	1.25	ANXA2.2	ANXA2	<0.0001	1.71
LGALS1.1	LGALS1	<0.0001	1.46	ALOX5.1	ALOX5	0.0004	1.38
LAPTM5.1	LAPTM5	<0.0001	1.47	ADAM8.1	ADAM8	<0.0001	1.89
LAMB3.1	LAMB3	<0.0001	1.96	MRP2.3	ABCC2	0.0002	1.39
L1CAM.1	L1CAM	<0.0001	1.43				

Table 6b: Proxy genes for which increased expression is associated with the absence of necrosis (p-value $\leq .05$)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
YB-1.2	YBX1	0.0010	0.74	IGF2.2	IGF2	0.0117	0.79
XIAP.1	XIAP	<0.0001	0.68	IGF1R.3	IGF1R	<0.0001	0.38
WWOX.5	WWOX	<0.0001	0.63	ID3.1	ID3	<0.0001	0.44
WISP1.1	WISP1	0.0002	0.71	ID2.4	ID2	<0.0001	0.68
VWF.1	VWF	<0.0001	0.35	ID1.1	ID1	<0.0001	0.32
VHL.1	VHL	0.0086	0.78	ICAM2.1	ICAM2	<0.0001	0.47
VEGF.1	VEGFA	<0.0001	0.50	HYAL2.1	HYAL2	<0.0001	0.28
VCAM1.1	VCAM1	<0.0001	0.55	HYAL1.1	HYAL1	<0.0001	0.39
USP34.1	USP34	<0.0001	0.64	HSPG2.1	HSPG2	<0.0001	0.33
UMOD.1	UMOD	<0.0001	0.36	HSP90AB1.1	HSP90AB1	0.0004	0.73
UGCG.1	UGCG	<0.0001	0.54	HSD11B2.1	HSD11B2	<0.0001	0.32
UBB.1	UBB	<0.0001	0.48	Hepsin.1	HPN	<0.0001	0.59
UBE1C.1	UBA3	<0.0001	0.59	HPCAL1.1	HPCAL1	<0.0001	0.68
TS.1	TYMS	<0.0001	0.70	HMGB1.1	HMGB1	<0.0001	0.42
tusc4.2	TUSC4	<0.0001	0.68	HLA-DPB1.1	HLA-DPB1	0.0002	0.72
TUSC2.1	TUSC2	0.0462	0.83	HIF1AN.1	HIF1AN	<0.0001	0.54
TSPAN7.2	TSPAN7	<0.0001	0.25	HDAC1.1	HDAC1	<0.0001	0.55
TSC2.1	TSC2	<0.0001	0.45	HAVCR1.1	HAVCR1	0.0012	0.76
TSC1.1	TSC1	<0.0001	0.45	HADH.1	HADH	<0.0001	0.49
P53.2	TP53	<0.0001	0.61	GSTT1.3	GSTT1	0.0067	0.80
TOP2B.2	TOP2B	<0.0001	0.69	GSTp.3	GSTP1	<0.0001	0.55
TNFSF12.1	TNFSF12	<0.0001	0.51	GSTM3.2	GSTM3	<0.0001	0.48
TRAIL.1	TNFSF10	<0.0001	0.68	GSTM1.1	GSTM1	<0.0001	0.54
TNFRSF11B.1	TNFRSF11B	<0.0001	0.59	GRB7.2	GRB7	<0.0001	0.49
TNFRSF10D.1	TNFRSF10D	<0.0001	0.58	GPX3.1	GPX3	<0.0001	0.59
DR5.2	TNFRSF10B	0.0001	0.71	GPC3.1	GPC3	0.0287	0.81
TNFAIP6.1	TNFAIP6	<0.0001	0.67	GJA1.1	GJA1	0.0004	0.74
TMEM47.1	TMEM47	<0.0001	0.29	GFRA1.1	GFRA1	0.0011	0.74
TMEM27.1	TMEM27	<0.0001	0.40	GCLC.3	GCLC	<0.0001	0.50
TLR3.1	TLR3	<0.0001	0.64	GATM.1	GATM	<0.0001	0.45
TIMP3.3	TIMP3	<0.0001	0.23	GATA3.3	GATA3	0.0159	0.79
TIMP2.1	TIMP2	<0.0001	0.52	GADD45B.1	GADD45B	<0.0001	0.67
THBS1.1	THBS1	<0.0001	0.62	FOS.1	FOS	<0.0001	0.56
TGFBR2.3	TGFBR2	<0.0001	0.34	FOLR1.1	FOLR1	<0.0001	0.59
TGFBR1.1	TGFBR1	<0.0001	0.63	FLT4.1	FLT4	<0.0001	0.27
TGFB2.2	TGFB2	<0.0001	0.55	FLT3LG.1	FLT3LG	<0.0001	0.62

Table 6b: Proxy genes for which increased expression is associated with the absence of necrosis (p-value $\leq .05$)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
TGFb1.1	TGFB1	0.0036	0.76	FLT1.1	FLT1	<0.0001	0.32
TGFA.2	TGFA	<0.0001	0.56	FILIP1.1	FILIP1	<0.0001	0.42
TEK.1	TEK	<0.0001	0.23	FIGF.1	FIGF	0.0001	0.62
TCF4.1	TCF4	<0.0001	0.36	FHL1.1	FHL1	<0.0001	0.36
TAGLN.1	TAGLN	<0.0001	0.50	FHIT.1	FHIT	<0.0001	0.63
TACSTD2.1	TACSTD2	<0.0001	0.64	FH.1	FH	<0.0001	0.65
SUCLG1.1	SUCLG1	<0.0001	0.50	FGFR2 isoform 1.1	FGFR2	<0.0001	0.51
STK11.1	STK11	<0.0001	0.48	FGFR1.3	FGFR1	<0.0001	0.55
STAT5B.2	STAT5B	<0.0001	0.36	FGF2.2	FGF2	<0.0001	0.61
STAT5A.1	STAT5A	<0.0001	0.56	FGF1.1	FGF1	<0.0001	0.49
STAT3.1	STAT3	<0.0001	0.63	FDPS.1	FDPS	<0.0001	0.43
SPRY1.1	SPRY1	<0.0001	0.42	FBXW7.1	FBXW7	<0.0001	0.60
SPAST.1	SPAST	0.0004	0.74	fas.1	FAS	0.0054	0.79
SPARCL1.1	SPARCL1	<0.0001	0.48	ESRRG.3	ESRRG	<0.0001	0.68
SPARC.1	SPARC	<0.0001	0.54	ERG.1	ERG	<0.0001	0.34
SOD1.1	SOD1	<0.0001	0.67	ERCC4.1	ERCC4	0.0197	0.81
SNRK.1	SNRK	<0.0001	0.25	ERCC1.2	ERCC1	<0.0001	0.60
SNAI1.1	SNAI1	0.0004	0.71	ERBB4.3	ERBB4	0.0037	0.72
MADH4.1	SMAD4	<0.0001	0.33	ErbB3.1	ERBB3	<0.0001	0.59
MADH2.1	SMAD2	<0.0001	0.40	HER2.3	ERBB2	<0.0001	0.40
SLC34A1.1	SLC34A1	<0.0001	0.47	EPHB4.1	EPHB4	<0.0001	0.44
SLC22A6.1	SLC22A6	<0.0001	0.52	EPHA2.1	EPHA2	<0.0001	0.36
SKIL.1	SKIL	<0.0001	0.59	EPAS1.1	EPAS1	<0.0001	0.29
SHANK3.1	SHANK3	<0.0001	0.25	ENPP2.1	ENPP2	<0.0001	0.50
SGK.1	SGK1	<0.0001	0.54	ENPEP.1	ENPEP	<0.0001	0.56
FRP1.3	SFRP1	0.0053	0.77	CD105.1	ENG	<0.0001	0.31
SEMA3F.3	SEMA3F	<0.0001	0.43	EMP1.1	EMP1	<0.0001	0.49
SELENBP1.1	SELENBP1	<0.0001	0.62	EMCN.1	EMCN	<0.0001	0.23
SDPR.1	SDPR	<0.0001	0.28	ELTD1.1	ELTD1	<0.0001	0.59
SDHA.1	SDHA	<0.0001	0.47	EIF2C1.1	EIF2C1	<0.0001	0.52
SCNN1A.2	SCNN1A	0.0013	0.73	EGR1.1	EGR1	<0.0001	0.54
SCN4B.1	SCN4B	<0.0001	0.35	EGLN3.1	EGLN3	<0.0001	0.69
S100A2.1	S100A2	0.0355	0.82	EGFR.2	EGFR	<0.0001	0.70
KIAA1303 raptor.1	RPTOR	<0.0001	0.49	EFNB2.1	EFNB2	<0.0001	0.34
RPS6KB1.3	RPS6KB1	<0.0001	0.55	EFNB1.2	EFNB1	<0.0001	0.41

Table 6b: Proxy genes for which increased expression is associated with the absence of necrosis (p-value $\leq .05$)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
RPS6KAI.1	RPS6KA1	0.0002	0.68	EEF1A1.1	EEF1A1	<0.0001	0.32
RPS23.1	RPS23	<0.0001	0.47	EDNRB.1	EDNRB	<0.0001	0.31
ROCK2.1	ROCK2	<0.0001	0.39	EDN2.1	EDN2	<0.0001	0.51
ROCK1.1	ROCK1	<0.0001	0.35	EDN1 endothelin.1	EDN1	<0.0001	0.41
RIPK1.1	RIPK1	<0.0001	0.45	EBAG9.1	EBAG9	0.0007	0.74
rhoC.1	RHOC	0.0001	0.70	DUSP1.1	DUSP1	<0.0001	0.65
RhoB.1	RHOB	<0.0001	0.41	DPYS.1	DPYS	<0.0001	0.66
ARHA.1	RHOA	<0.0001	0.45	DPEP1.1	DPEP1	<0.0001	0.34
RHEB.2	RHEB	0.0002	0.69	DLL4.1	DLL4	<0.0001	0.49
RGS5.1	RGS5	<0.0001	0.26	DLC1.1	DLC1	<0.0001	0.36
FLJ22655.1	RERGL	<0.0001	0.26	DKFZP564O08 23.1	DKFZP564O 0823	<0.0001	0.35
NFKBp65.3	RELA	<0.0001	0.71	DICER1.2	DICER1	<0.0001	0.46
RB1.1	RB1	<0.0001	0.54	DIAPH1.1	DIAPH1	<0.0001	0.63
RASSF1.1	RASSF1	<0.0001	0.63	DIABLO.1	DIABLO	0.0002	0.72
RARB.2	RARB	<0.0001	0.28	DHPS.3	DHPS	<0.0001	0.46
RALBP1.1	RALBP1	<0.0001	0.52	DET1.1	DET1	<0.0001	0.61
RAF1.3	RAF1	<0.0001	0.58	DEFB1.1	DEFB1	<0.0001	0.69
RAC1.3	RAC1	0.0118	0.80	DDC.1	DDC	<0.0001	0.51
PTPRG.1	PTPRG	<0.0001	0.27	DCXR.1	DCXR	0.0061	0.78
PTPRB.1	PTPRB	<0.0001	0.22	DAPK1.3	DAPK1	<0.0001	0.47
PTN.1	PTN	<0.0001	0.41	CYR61.1	CYR61	<0.0001	0.52
PTK2.1	PTK2	<0.0001	0.33	CYP3A4.2	CYP3A4	0.0398	0.82
PTHR1.1	PTH1R	<0.0001	0.32	CYP2C8v2.1	CYP2C8_21	0.0001	0.67
PTEN.2	PTEN	<0.0001	0.44	CXCL12.1	CXCL12	<0.0001	0.53
PSMA7.1	PSMA7	0.0173	0.81	CX3CR1.1	CX3CR1	<0.0001	0.60
PRSS8.1	PRSS8	<0.0001	0.62	CX3CL1.1	CX3CL1	<0.0001	0.34
PRKCH.1	PRKCH	<0.0001	0.43	CUL1.1	CUL1	<0.0001	0.62
PRKCD.2	PRKCD	<0.0001	0.68	CUBN.1	CUBN	<0.0001	0.39
PPP2CA.1	PPP2CA	<0.0001	0.53	CTSH.2	CTSH	0.0018	0.77
PPARG.3	PPARG	<0.0001	0.37	B-Catenin.3	CTNNB1	<0.0001	0.38
PPAP2B.1	PPAP2B	<0.0001	0.27	A-Catenin.2	CTNNA1	<0.0001	0.58
PMP22.1	PMP22	0.0155	0.81	CTGF.1	CTGF	<0.0001	0.64
PLG.1	PLG	<0.0001	0.38	CSF1R.2	CSF1R	0.0308	0.83
PLAT.1	PLAT	<0.0001	0.42	CSF1.1	CSF1	0.0002	0.73
PLA2G4C.1	PLA2G4C	<0.0001	0.48	CRADD.1	CRADD	<0.0001	0.60

Table 6b: Proxy genes for which increased expression is associated with the absence of necrosis (p-value $\leq$.05)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
PIK3CA.1	PIK3CA	<0.0001	0.47	COL4A2.1	COL4A2	<0.0001	0.51
PI3K.2	PIK3C2B	<0.0001	0.42	COL4A1.1	COL4A1	<0.0001	0.66
PGF.1	PGF	0.0153	0.80	COL18A1.1	COL18A1	<0.0001	0.50
PFKP.1	PFKP	<0.0001	0.70	CLU.3	CLU	0.0091	0.80
CD31.3	PECAM1	<0.0001	0.34	CLDN7.2	CLDN7	0.0029	0.76
PDZK3.1	PDZK3	<0.0001	0.49	CLDN10.1	CLDN10	<0.0001	0.48
PDZK1.1	PDZK1	<0.0001	0.45	CLCNKB.1	CLCNKB	0.0001	0.61
PDGFRb.3	PDGFRB	<0.0001	0.45	CFLAR.1	CFLAR	<0.0001	0.47
PDGFD.2	PDGFD	<0.0001	0.33	CEACAM1.1	CEACAM1	<0.0001	0.43
PDGFC.3	PDGFC	<0.0001	0.53	p27.3	CDKN1B	0.0002	0.73
PDGFB.3	PDGFB	<0.0001	0.33	p21.3	CDKN1A	<0.0001	0.65
PDGFA.3	PDGFA	<0.0001	0.43	CDH6.1	CDH6	<0.0001	0.67
PCK1.1	PCK1	<0.0001	0.44	CDH5.1	CDH5	<0.0001	0.33
PCCA.1	PCCA	<0.0001	0.47	CDH2.1	CDH2	0.0003	0.75
PARD6A.1	PARD6A	0.0045	0.77	CDH16.1	CDH16	<0.0001	0.51
Pak1.2	PAK1	0.0003	0.74	CDH13.1	CDH13	<0.0001	0.39
PAH.1	PAH	<0.0001	0.62	CD36.1	CD36	<0.0001	0.41
OGG1.1	OGG1	<0.0001	0.62	CD34.1	CD34	<0.0001	0.34
BFGF.3	NUDT6	<0.0001	0.45	CD24.1	CD24	0.0148	0.81
NRG1.3	NRG1	0.0004	0.69	CCND1.3	CCND1	<0.0001	0.51
NPR1.1	NPR1	<0.0001	0.36	MCP1.1	CCL2	<0.0001	0.68
NPM1.2	NPM1	<0.0001	0.55	CAT.1	CAT	<0.0001	0.48
NOTCH3.1	NOTCH3	<0.0001	0.40	CASP10.1	CASP10	<0.0001	0.62
NOTCH2.1	NOTCH2	<0.0001	0.68	CALD1.2	CALD1	<0.0001	0.42
NOTCH1.1	NOTCH1	<0.0001	0.38	CACNA2D1.1	CACNA2D1	0.0006	0.74
NOS3.1	NOS3	<0.0001	0.37	CA2.1	CA2	<0.0001	0.60
NOS2A.3	NOS2	<0.0001	0.42	C7.1	C7	<0.0001	0.65
NOL3.1	NOL3	<0.0001	0.67	ECRG4.1	C2orf40	<0.0001	0.32
NFX1.1	NFX1	<0.0001	0.43	C13orf15.1	C13orf15	<0.0001	0.31
NFKBp50.3	NFKB1	<0.0001	0.56	BUB3.1	BUB3	<0.0001	0.65
NFATC2.1	NFATC2	<0.0001	0.67	BTRC.1	BTRC	<0.0001	0.63
NFAT5.1	NFAT5	<0.0001	0.55	BNIP3.1	BNIP3	0.0021	0.77
MYRIP.2	MYRIP	<0.0001	0.36	CIAP1.2	BIRC2	<0.0001	0.56
MYH11.1	MYH11	<0.0001	0.35	BIN1.3	BIN1	<0.0001	0.67
cMYC.3	MYC	<0.0001	0.68	BGN.1	BGN	<0.0001	0.47
MVP.1	MVP	<0.0001	0.66	BCL2L12.1	BCL2L12	0.0374	0.82

Table 6b: Proxy genes for which increased expression is associated with the absence of necrosis (p-value $\leq$.05)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
FRAP1.1	MTOR	<0.0001	0.56	Bclx.2	BCL2L1	<0.0001	0.60
MSH3.2	MSH3	<0.0001	0.47	Bcl2.2	BCL2	<0.0001	0.31
MSH2.3	MSH2	<0.0001	0.51	BAG1.2	BAG1	<0.0001	0.42
MMP2.2	MMP2	0.0229	0.82	BAD.1	BAD	0.0187	0.82
STMY3.3	MMP11	<0.0001	0.66	AXL.1	AXL	0.0077	0.79
GBL.1	MLST8	0.0193	0.82	ATP6V1B1.1	ATP6V1B1	<0.0001	0.52
MIF.2	MIF	<0.0001	0.69	ASS1.1	ASS1	<0.0001	0.61
MICA.1	MICA	<0.0001	0.52	ARRB1.1	ARRB1	<0.0001	0.48
MGMT.1	MGMT	<0.0001	0.50	ARHGDIB.1	ARHGDIB	<0.0001	0.48
MCM3.3	MCM3	0.0105	0.78	ARF1.1	ARF1	0.0021	0.75
MCAM.1	MCAM	<0.0001	0.41	AQP1.1	AQP1	<0.0001	0.28
MARCKS.1	MARCKS	0.0259	0.82	APOLD1.1	APOLD1	<0.0001	0.35
ERK1.3	MAPK3	<0.0001	0.35	APC.4	APC	<0.0001	0.55
ERK2.3	MAPK1	<0.0001	0.61	APAF1.2	APAF1	0.0264	0.82
MAP4.1	MAP4	<0.0001	0.54	ANXA4.1	ANXA4	0.0012	0.76
MAP2K3.1	MAP2K3	<0.0001	0.52	ANXA1.2	ANXA1	0.0201	0.81
MAP2K1.1	MAP2K1	0.0172	0.81	ANTXR1.1	ANTXR1	<0.0001	0.58
MAL2.1	MAL2	0.0267	0.82	ANGPTL4.1	ANGPTL4	<0.0001	0.71
MAL.1	MAL	<0.0001	0.46	ANGPTL3.3	ANGPTL3	0.0104	0.77
LTF.1	LTF	0.0038	0.74	ANGPTL2.1	ANGPTL2	<0.0001	0.63
LRP2.1	LRP2	<0.0001	0.52	ANGPT2.1	ANGPT2	<0.0001	0.65
LMO2.1	LMO2	<0.0001	0.60	ANGPT1.1	ANGPT1	<0.0001	0.30
LDB2.1	LDB2	<0.0001	0.26	AMACR1.1	AMACR	0.0080	0.79
LDB1.2	LDB1	<0.0001	0.52	ALDOB.1	ALDOB	<0.0001	0.40
LAMA4.1	LAMA4	<0.0001	0.67	ALDH6A1.1	ALDH6A1	<0.0001	0.38
KRT7.1	KRT7	<0.0001	0.56	ALDH4.2	ALDH4A1	0.0001	0.71
K-ras.10	KRAS	<0.0001	0.48	AKT3.2	AKT3	<0.0001	0.30
KL.1	KL	<0.0001	0.34	AKT2.3	AKT2	<0.0001	0.53
Kitlng.4	KITLG	<0.0001	0.46	AKT1.3	AKT1	<0.0001	0.47
c-kit.2	KIT	<0.0001	0.41	AHR.1	AHR	<0.0001	0.60
KDR.6	KDR	<0.0001	0.28	AGTR1.1	AGTR1	<0.0001	0.33
KCNJ15.1	KCNJ15	<0.0001	0.43	AGT.1	AGT	0.0032	0.77
HTATIP.1	KAT5	<0.0001	0.27	ADH6.1	ADH6	0.0011	0.71
G-Catenin.1	JUP	<0.0001	0.32	ADH1B.1	ADH1B	<0.0001	0.69
AP-1 (JUN official).2	JUN	<0.0001	0.64	ADFP.1	ADFP	0.0001	0.73

Table 6b: Proxy genes for which increased expression is associated with the absence of necrosis (p-value $\leq$.05)

Gene	Official Symbol	CCF Necrosis		Gene	Official Symbol	CCF Necrosis	
		p-value	OR			p-value	OR
JAG1.1	JAG1	<0.0001	0.23	ADD1.1	ADD1	<0.0001	0.33
ITGB5.1	ITGB5	<0.0001	0.64	ADAMTS9.1	ADAMTS9	<0.0001	0.69
ITGB3.1	ITGB3	0.0468	0.84	ADAMTS5.1	ADAMTS5	<0.0001	0.55
ITGB1.1	ITGB1	<0.0001	0.59	ADAMTS1.1	ADAMTS1	<0.0001	0.56
ITGA7.1	ITGA7	<0.0001	0.38	ADAM17.1	ADAM17	0.0009	0.76
ITGA6.2	ITGA6	<0.0001	0.40	ACE2.1	ACE2	<0.0001	0.45
ITGA5.1	ITGA5	0.0298	0.83	ACADSB.1	ACADSB	<0.0001	0.46
ITGA4.2	ITGA4	<0.0001	0.61	BCRP.1	ABCG2	<0.0001	0.27
ITGA3.2	ITGA3	0.0018	0.76	MRP4.2	ABCC4	<0.0001	0.61
IQGAP2.1	IQGAP2	<0.0001	0.52	MRP3.1	ABCC3	0.0011	0.76
INSR.1	INSR	<0.0001	0.38	MRP1.1	ABCC1	0.0008	0.75
IMP3.1	IMP3	<0.0001	0.53	ABCB1.5	ABCB1	<0.0001	0.59
IL6ST.3	IL6ST	<0.0001	0.36	NPD009 (ABAT official).3	ABAT	0.0001	0.70
IL15.1	IL15	<0.0001	0.67	AAMP.1	AAMP	<0.0001	0.62
IGFBP6.1	IGFBP6	<0.0001	0.66	A2M.1	A2M	<0.0001	0.28

Table 7a: Proxy genes for which increased expression is associated with the presence of nodal invasion (p-value $\leq$.05)

Gene	Official Symbol	Nodal Invasion		Gene	Official Symbol	Nodal Invasion	
		p-value	OR			p-value	OR
TUBB.1	TUBB2A	0.0242	2.56	IL-8.1	IL8	0.0019	3.18
C20 orf1.1	TPX2	0.0333	2.61	IL6.3	IL6	0.0333	2.31
TK1.2	TK1	0.0361	1.75	HSPA1A.1	HSPA1A	0.0498	2.11
SPHK1.1	SPHK1	0.0038	3.43	GSTp.3	GSTP1	0.0272	3.46
SLC7A5.2	SLC7A5	0.0053	4.85	GRB14.1	GRB14	0.0287	2.32
SILV.1	SILV	0.0470	1.54	GMNN.1	GMNN	0.0282	3.00
SELE.1	SELE	0.0311	1.93	ENO2.1	ENO2	0.0190	3.43
upa.3	PLAU	0.0450	2.78	CCNB1.2	CCNB1	0.0387	1.87
MMP9.1	MMP9	0.0110	2.65	BUB1.1	BUB1	0.0429	2.21
MMP7.1	MMP7	0.0491	2.34	BAG2.1	BAG2	0.0346	2.54
MMP14.1	MMP14	0.0155	3.21	ADAMTS1.1	ADAMTS1	0.0193	3.31
LAMB1.1	LAMB1	0.0247	3.04				

Table 7b: Proxy genes for which increased expression is associated with the absence of nodal invasion (p-value $\leq .05$)

Gene	Official Symbol	Nodal Invasion		Gene	Official Symbol	Nodal Invasion	
		p-value	OR			p-value	OR
VWF.1	VWF	0.0221	0.42	HMGB1.1	HMGB1	0.0385	0.45
VCAM1.1	VCAM1	0.0212	0.42	HLA-DPB1.1	HLA-DPB1	0.0398	0.43
UBE1C.1	UBA3	0.0082	0.32	HADH.1	HADH	0.0093	0.33
tusc4.2	TUSC4	0.0050	0.28	GSTM1.1	GSTM1	0.0018	0.18
TSPAN7.2	TSPAN7	0.0407	0.43	GPX2.2	GPX2	0.0211	0.07
TSC1.1	TSC1	0.0372	0.38	GJA1.1	GJA1	0.0451	0.46
TMSB10.1	TMSB10	0.0202	0.43	GATM.1	GATM	0.0038	0.25
TMEM47.1	TMEM47	0.0077	0.32	GATA3.3	GATA3	0.0188	0.06
TMEM27.1	TMEM27	0.0431	0.41	FOLR1.1	FOLR1	0.0152	0.32
TLR3.1	TLR3	0.0041	0.32	FLT4.1	FLT4	0.0125	0.22
TIMP3.3	TIMP3	0.0309	0.40	FLT1.1	FLT1	0.0046	0.35
TGFBR2.3	TGFBR2	0.0296	0.35	FHL1.1	FHL1	0.0435	0.47
TGFB2.2	TGFB2	0.0371	0.30	FHIT.1	FHIT	0.0061	0.29
TGFA.2	TGFA	0.0025	0.32	fas.1	FAS	0.0163	0.39
TEK.1	TEK	0.0018	0.09	ErbB3.1	ERBB3	0.0145	0.33
TCF4.1	TCF4	0.0088	0.38	EPHA2.1	EPHA2	0.0392	0.37
STAT5A.1	STAT5A	0.0129	0.49	EPAS1.1	EPAS1	0.0020	0.28
SPRY1.1	SPRY1	0.0188	0.43	ENPEP.1	ENPEP	0.0002	0.34
SPARCL1.1	SPARCL1	0.0417	0.50	CD105.1	ENG	0.0112	0.38
SOD1.1	SOD1	0.0014	0.23	EMCN.1	EMCN	0.0022	0.19
SNRK.1	SNRK	0.0226	0.43	EIF2C1.1	EIF2C1	0.0207	0.33
MADH2.1	SMAD2	0.0098	0.37	EGLN3.1	EGLN3	0.0167	0.52
SLC22A6.1	SLC22A6	0.0051	0.00	EFNB2.1	EFNB2	0.0192	0.30
PTPNS1.1	SIRPA	0.0206	0.43	EFNB1.2	EFNB1	0.0110	0.37
SHANK3.1	SHANK3	0.0024	0.30	EDNRB.1	EDNRB	0.0106	0.36
SGK.1	SGK1	0.0087	0.35	EDN1 endothelin.1	EDN1	0.0440	0.42
SELENBP1.1	SELENBP1	0.0016	0.29	DPYS.1	DPYS	0.0454	0.43
SCN4B.1	SCN4B	0.0081	0.08	DKFZP564O08 23.1	DKFZP564O 0823	0.0131	0.31
ROCK1.1	ROCK1	0.0058	0.41	DHPS.3	DHPS	0.0423	0.48
RhoB.1	RHOB	0.0333	0.43	DAPK1.3	DAPK1	0.0048	0.39
RGS5.1	RGS5	0.0021	0.31	CYP2C8v2.1	CYP2C8_21	0.0003	0.01
FLJ22655.1	RERGL	0.0009	0.01	CYP2C8.2	CYP2C8_2	0.0269	0.07
RB1.1	RB1	0.0281	0.48	CX3CR1.1	CX3CR1	0.0224	0.13
RASSF1.1	RASSF1	0.0004	0.23	CUBN.1	CUBN	0.0010	0.04

Table 7b: Proxy genes for which increased expression is associated with the absence of nodal invasion (p-value $\leq .05$)

Gene	Official Symbol	Nodal Invasion		Gene	Official Symbol	Nodal Invasion	
		p-value	OR			p-value	OR
PTPRB.1	PTPRB	0.0154	0.40	CRADD.1	CRADD	0.0193	0.40
PTK2.1	PTK2	0.0158	0.38	CLDN10.1	CLDN10	0.0005	0.21
PTHR1.1	PTH1R	<0.0001	0.01	CFLAR.1	CFLAR	0.0426	0.49
PRSS8.1	PRSS8	0.0023	0.10	CEACAM1.1	CEACAM1	0.0083	0.23
PRKCH.1	PRKCH	0.0475	0.48	CDKN2A.2	CDKN2A	0.0026	0.13
PPAP2B.1	PPAP2B	0.0110	0.40	p27.3	CDKN1B	0.0393	0.60
PLA2G4C.1	PLA2G4C	0.0002	0.03	CDH5.1	CDH5	0.0038	0.33
PI3K.2	PIK3C2B	0.0138	0.13	CDH13.1	CDH13	0.0203	0.39
PFKP.1	PFKP	0.0040	0.41	CD99.1	CD99	0.0173	0.38
CD31.3	PECAM1	0.0077	0.38	CD36.1	CD36	0.0015	0.29
PDGFD.2	PDGFD	0.0169	0.35	CD34.1	CD34	0.0250	0.38
PDGFC.3	PDGFC	0.0053	0.36	CD3z.1	CD247	0.0419	0.33
PDGFB.3	PDGFB	0.0359	0.47	CCND1.3	CCND1	0.0381	0.50
PCSK6.1	PCSK6	0.0103	0.34	CAT.1	CAT	0.0044	0.42
PCK1.1	PCK1	0.0003	0.02	CASP6.1	CASP6	0.0136	0.36
PCCA.1	PCCA	0.0074	0.12	CALD1.2	CALD1	0.0042	0.31
PARD6A.1	PARD6A	0.0243	0.21	CA9.3	CA9	0.0077	0.48
BFGF.3	NUDT6	0.0082	0.15	C13orf15.1	C13orf15	0.0152	0.39
NRG1.3	NRG1	0.0393	0.19	BUB3.1	BUB3	0.0157	0.26
NOS3.1	NOS3	0.0232	0.34	BIN1.3	BIN1	0.0224	0.38
NOS2A.3	NOS2	0.0086	0.11	Bclx.2	BCL2L1	0.0225	0.52
NFX1.1	NFX1	0.0065	0.35	Bcl2.2	BCL2	0.0442	0.45
MYH11.1	MYH11	0.0149	0.36	AXL.1	AXL	0.0451	0.44
cMYC.3	MYC	0.0472	0.46	ATP6V1B1.1	ATP6V1B1	0.0257	0.03
MUC1.2	MUC1	0.0202	0.26	ARRB1.1	ARRB1	0.0337	0.43
MIF.2	MIF	0.0145	0.39	ARHGDIB.1	ARHGDIB	0.0051	0.32
MICA.1	MICA	0.0015	0.15	AQP1.1	AQP1	0.0022	0.31
MGMT.1	MGMT	0.0414	0.50	APOLD1.1	APOLD1	0.0222	0.42
MAP2K1.1	MAP2K1	0.0075	0.40	APC.4	APC	0.0165	0.47
LMO2.1	LMO2	0.0127	0.07	ANXA5.1	ANXA5	0.0143	0.37
LDB2.1	LDB2	0.0048	0.33	ANXA4.1	ANXA4	0.0019	0.30
Kitlng.4	KITLG	0.0146	0.18	ANXA1.2	ANXA1	0.0497	0.39
KDR.6	KDR	0.0106	0.34	ANGPTL7.1	ANGPTL7	0.0444	0.16
ITGB1.1	ITGB1	0.0469	0.36	ANGPTL4.1	ANGPTL4	0.0197	0.55
ITGA7.1	ITGA7	0.0290	0.38	ANGPT1.1	ANGPT1	0.0055	0.15

Table 7b: Proxy genes for which increased expression is associated with the absence of nodal invasion (p-value $\leq$.05)

Gene	Official Symbol	Nodal Invasion		Gene	Official Symbol	Nodal Invasion	
		p-value	OR			p-value	OR
ITGA6.2	ITGA6	0.0010	0.17	ALDOB.1	ALDOB	0.0128	0.02
ITGA4.2	ITGA4	0.0089	0.44	ALDH4.2	ALDH4A1	0.0304	0.33
INSR.1	INSR	0.0057	0.32	AGTR1.1	AGTR1	0.0142	0.11
IMP3.1	IMP3	0.0086	0.43	ADH6.1	ADH6	0.0042	0.03
IL6ST.3	IL6ST	0.0484	0.46	ADFP.1	ADFP	0.0223	0.47
IL15.1	IL15	0.0009	0.08	ADD1.1	ADD1	0.0135	0.42
IFI27.1	IFI27	0.0013	0.22	BCRP.1	ABCG2	0.0098	0.19
HYAL2.1	HYAL2	0.0099	0.36	MRP3.1	ABCC3	0.0247	0.44
HYAL1.1	HYAL1	0.0001	0.08	MRP1.1	ABCC1	0.0022	0.32
Hepsin.1	HPN	0.0024	0.14	NPD009 (ABAT official).3	ABAT	0.0110	0.19
HPCAL1.1	HPCAL1	0.0024	0.38	A2M.1	A2M	0.0012	0.29

Table 8a: Genes for which increased expression is associated with lower risk of cancer recurrence after clinical/pathologic covariate adjustment (p<0.05)

Gene	Official Symbol	p-value	HR	Gene	Official Symbol	p-value	HR
ACE2.1	ACE2	0.0261	0.85	ID1.1	ID1	0.0154	0.83
ADD1.1	ADD1	0.0339	0.85	IGF1R.3	IGF1R	0.0281	0.85
ALDOB.1	ALDOB	0.0328	0.84	IL15.1	IL15	0.0059	0.83
ANGPTL3.3	ANGPTL3	0.0035	0.79	IQGAP2.1	IQGAP2	0.0497	0.88
APOLD1.1	APOLD1	0.0015	0.78	KL.1	KL	0.0231	0.86
AQP1.1	AQP1	0.0014	0.79	KLRK1.2	KLRK1	0.0378	0.87
BFGF.3	NUDT6	0.0010	0.77	LDB2.1	LDB2	0.0092	0.82
CASP10.1	CASP10	0.0024	0.82	LRP2.1	LRP2	0.0193	0.86
CAV2.1	CAV2	0.0191	0.86	LTF.1	LTF	0.0077	0.82
CCL4.2	CCL4	0.0045	0.81	MAP4.1	MAP4	0.0219	0.84
CCL5.2	CCL5	0.0003	0.78	MRP1.1	ABCC1	0.0291	0.87
CCR2.1	CCR2	0.0390	0.87	NOS3.1	NOS3	0.0008	0.78
CCR4.2	CCR4	0.0109	0.82	PI3K.2	PIK3C2B	0.0329	0.83
CCR7.1	CCR7	0.0020	0.80	PLA2G4C.1	PLA2G4C	0.0452	0.87
CD4.1	CD4	0.0195	0.86	PPAP2B.1	PPAP2B	0.0001	0.74
CD8A.1	CD8A	0.0058	0.83	PRCC.1	PRCC	0.0333	0.88
CEACAM1.1	CEACAM1	0.0022	0.81	PRKCB1.1	PRKCB	0.0353	0.87
CFLAR.1	CFLAR	0.0308	0.87	PRKCH.1	PRKCH	0.0022	0.82
CTSS.1	CTSS	0.0462	0.87	PRSS8.1	PRSS8	0.0332	0.87
CX3CL1.1	CX3CL1	0.0021	0.81	PSMB9.1	PSMB9	0.0262	0.87
CXCL10.1	CXCL10	0.0323	0.86	PTPRB.1	PTPRB	0.0030	0.80
CXCL9.1	CXCL9	0.0006	0.79	RGS5.1	RGS5	0.0480	0.85
CXCR6.1	CXCR6	0.0469	0.88	SDPR.1	SDPR	0.0045	0.80
DAPK1.3	DAPK1	0.0050	0.83	SELE.1	SELE	0.0070	0.81
DDC.1	DDC	0.0307	0.86	SGK.1	SGK1	0.0100	0.83
DLC1.1	DLC1	0.0249	0.83	SHANK3.1	SHANK3	0.0311	0.84
ECRG4.1	C2orf40	0.0244	0.84	SNRK.1	SNRK	0.0026	0.81
EDNRB.1	EDNRB	0.0400	0.86	TEK.1	TEK	0.0059	0.78
EMCN.1	EMCN	<0.0001	0.68	TGFBR2.3	TGFBR2	0.0343	0.85
EPAS1.1	EPAS1	0.0411	0.84	TIMP3.3	TIMP3	0.0165	0.83
fas.1	FAS	0.0242	0.87	TMEM27.1	TMEM27	0.0249	0.86
FH.1	FH	0.0407	0.88	TSPAN7.2	TSPAN7	0.0099	0.83
GATA3.3	GATA3	0.0172	0.83	UBB.1	UBB	0.0144	0.85
GZMA.1	GZMA	0.0108	0.84	WWOX.5	WWOX	0.0082	0.83
HLA-DPB1.1	HLA-DPB1	0.0036	0.82				
HSPG2.1	HSPG2	0.0236	0.84				
ICAM2.1	ICAM2	0.0091	0.83				
ICAM3.1	ICAM3	0.0338	0.87				

Table 8b: Genes for which increased expression is associated with higher risk of cancer recurrence after clinical/pathologic covariate adjustment ($p < 0.05$)

Gene	Official Symbol	p-value	HR
CIAP1.2	BIRC2	0.0425	1.14
BUB1.1	BUB1	0.0335	1.15
CCNB1.2	CCNB1	0.0296	1.14
ENO2.1	ENO2	0.0284	1.17
ITGB1.1	ITGB1	0.0402	1.16
ITGB5.1	ITGB5	0.0016	1.25
LAMB1.1	LAMB1	0.0067	1.20
MMP14.1	MMP14	0.0269	1.16
MMP9.1	MMP9	0.0085	1.19
PSMA7.1	PSMA7	0.0167	1.16
RUNX1.1	RUNX1	0.0491	1.15
SPHK1.1	SPHK1	0.0278	1.16
OPN, osteopontin.3	SPP1	0.0134	1.17
SQSTM1.1	SQSTM1	0.0347	1.13
C20 orf1.1	TPX2	0.0069	1.20
TUBB.1	TUBB2A	0.0046	1.21
VCAN.1	VCAN	0.0152	1.18

Table 9: 16 Significant Genes After Adjusting for Clinical/Pathologic Covariates and Allowing for an FDR of 10%

Official Symbol	n	Gene Subset (Pathway)	HR	HR (95% CI)	LR ChiSq	LR p-value	q-value
EMCN	928	Angiogenesis	0.68	(0.57,0.80)	19.87	<0.0001	0.0042
PPAP2B	928	Angiogenesis	0.74	(0.65,0.85)	16.15	0.0001	0.0148
CCL5	928	Immune Response	0.78	(0.68,0.89)	12.98	0.0003	0.0529
CXCL9	928	Immune Response	0.79	(0.70,0.91)	11.74	0.0006	0.0772
NOS3	928	Angiogenesis	0.78	(0.68,0.90)	11.32	0.0008	0.0774
NUDT6	926	Angiogenesis	0.77	(0.66,0.90)	10.77	0.0010	0.0850
AQP1	928	Transport	0.79	(0.69,0.91)	10.15	0.0014	0.0850
APOLD1	927	Angiogenesis	0.78	(0.68,0.91)	10.07	0.0015	0.0850
ITGB5	928	Cell Adhesion/ Extracellular Matrix	1.25	(1.09,1.43)	9.92	0.0016	0.0850
CCR7	928	Immune Response	0.80	(0.69,0.92)	9.58	0.0020	0.0850
CX3CL1	926	Immune Response	0.81	(0.70,0.92)	9.44	0.0021	0.0850
CEACAM1	928	Angiogenesis	0.81	(0.70,0.93)	9.37	0.0022	0.0850
PRKCH	917	Angiogenesis	0.82	(0.72,0.93)	9.36	0.0022	0.0850
CASP10	927	Apoptosis	0.82	(0.73,0.93)	9.25	0.0024	0.0850
SNRK	928	Angiogenesis	0.81	(0.71,0.92)	9.09	0.0026	0.0863
PTPRB	927	Angiogenesis	0.80	(0.69,0.92)	8.83	0.0030	0.0933

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
A-Catenin.2	NM_001903	NM_001903.1	CTNNA1	765	CGTTCCGATCCTCT ATACTGCAT	AGGTCCCTGTTG GCCTTATAGG	ATGCCTACAGCACCC TGATGTCGCA
A2M.1	NM_000014	NM_000014.4	A2M	6456	CTCTCCCGCCTTCC TAGC	CCGTTTGCACAGA TGCAG	CGCTTGTTCTTCTC CACTGGGAC
AAMP.1	NM_001087	NM_001087.3	AAMP	5474	GTGTGGCAGGTGG ACACTAA	CTCCATCCACTCC AGGTCTC	CGCTTCAAAGGACCA GACCTCCTC
ABCB1.5	NM_000927	NM_000927.2	ABCB1	3099	AAACACCACTGGAG CATTGA	CAAGCCTGGAAC CTATAGCC	CTCGCCAATGATGCT GCTCAAGTT
ACADSB.1	NM_001609	NM_001609.3	ACADSB	6278	TGGCGGAGAACTA GCCAT	AAGACAGCCCAG TCCTCAAAT	CCTCCTGAAGCCTG CCATCATTGT
ACE.1	NM_000789	NM_000789.2	ACE	4257	CCGCTGTACGAGG ATTTC	CCGTGTCTGTGA AGCCGT	TGCCCTCAGCAATGA AGCCTACAA
ACE2.1	NM_021804	NM_021804.1	ACE2	6108	TACAATGAGAGGCT CTGGGC	TAATGGCCTCAGC TGCTTG	CGACCTCAGATCTCC AGCTTTCCC
ADAM17.1	NM_003183	NM_003183.3	ADAM17	2617	GAAGTGCCAGGAG GCGATTA	CGGGCACTCACT GCTATTACC	TGCTACTTGCAAAGG CGTGTCTTACTGC
ADAM8.1	NM_001109	NM_001109.2	ADAM8	3978	GTCCTGTGTCCAG CCCA	TGATGACCTGCTT TGGTGC	TTCCCAGTTCTGTG TACACCCGG
ADAMTS1.1	NM_006988	NM_006988.2	ADAMTS1	2639	GGACAGGTGCAAG CTCATCTG	ATCTACAACCTTG GGCTGCAA	CAAGCCAAAGGCATT GGCTACTTCTTCG
ADAMTS2.1	NM_014244	NM_014244.1	ADAMTS2	3979	GAGAATGTCTGCC GCTGG	ATCGTGCTATTCA TCGTGGC	TACCTCCAGCAGAAG CCAGACACG
ADAMTS4.1	NM_005099	NM_005099.3	ADAMTS4	2642	TTTGACAAGTGCAT GGTGTG	AATTCCTGAAGG AGCCTGA	CTGCTTGCTGCAACC AGAACCGT
ADAMTS5.1	NM_007038	NM_007038.1	ADAMTS5	2641	CACTGTGGCTCAC GAAATCG	GGAACCAAAGGT CTCTTCACAGA	ATTTACTTGGCCTCT CCCATGACGATTCC
ADAMTS8.1	NM_007037	NM_007037.2	ADAMTS8	2640	GCGAGTTCAAAGTG TTCGAG	CACAGATGGCCA GTGTTTCT	CACACAGGGTGCCA TCAATCACCT
ADAMTS9.1	NM_182920	NM_182920.1	ADAMTS9	6109	GCACAGGTTACACA ACCCAA	CGACATTGGCAG TCATCG	CCGGCTCCCGTTATA GGGACATTC
ADD1.1	NM_001119	NM_001119.3	ADD1	3980	GTCTACCCAGCAG CTCCG	TCGTTTACAGGA GTCACCAT	CATGTTTAAGGCAGC CATCCCTCC
ADFP.1	NM_001122	NM_001122.2	PLIN2	4503	AAGACCATCACCTC CGTGG	CAATTTGCGGCTC TAGCTTC	ATGACCAGTGCTCTG CCCATCATC
ADH1B.1	NM_000668	NM_000668.4	ADH1B	6325	AAGCCAACAAACCT TCCTTC	AAAATGCAAGAAG TCACAGGAA	TTTCCTCAATGGCAA AGGTGACACA

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
ADH6.1	NM_000672	NM_000672.3	ADH6	6111	TGTTGGGGAGTAAA CACTTGG	AACGATTCCAGCC CCTTC	TCTTGTATCCCACCA TCTTGGGCC
ADM.1	NM_001124	NM_001124.1	ADM	3248	TAAGCCACAAGCAC ACGG	TGGGCGCCTAAA TCCTAA	CGAGTGGAAGTGCT CCCCACTTTC
AGR2.1	NM_006408	NM_006408.2	AGR2	3245	AGCCAACATGTGAC TAATTGGA	TCTGATCTCCATC TGCCTCA	CAACACGTCACCACC CTTTGCTCT
AGT.1	NM_000029	NM_000029.2	AGT	6112	GATCCAGCCTCACT ATGCCT	CCAGTTGAGGGA GTTTTGCT	TGAGACCCTCCACCT TGTCCAGGT
AGTR1.1	NM_000685	NM_000685.3	AGTR1	4258	AGCATTGATCGATA CCTGGC	CTACAAGCATTGT GCGTCG	ATTGTTCACCCAATG AAGTCCCGC
AHR.1	NM_001621	NM_001621.2	AHR	3981	GCGGCATAGAGAC CGACTT	ACATCTTGTGGGA AAGGCA	CAGGCTAGCCAAAC GGTCCAACTC
AIF1.1	NM_032955	NM_032955.1	AIF1	6452	GACGTTTCAGCTACC CTGACTTT	TCAGGATCATTTT TAGGATGGC	ATCTCTTGCCAGCA TCATCCTGA
AKT1.3	NM_005163	NM_005163.1	AKT1	18	CGCTTCTATGGCGC TGAGAT	TCCCGGTACACC ACGTTCTT	CAGCCCTGGACTAC CTGCACTCGG
AKT2.3	NM_001626	NM_001626.2	AKT2	358	TCCTGCCACCCTTC AAACC	GGCGGTAAATTCA TCATCGAA	CAGGTCACGTCCGA GGTCGACACA
AKT3.2	NM_005465	NM_005465.1	AKT3	21	TTGTCTCTGCCTTG GACTATCTACA	CCAGCATTAGATT CTCCAACTTGA	TCACGGTACACAATC TTTCCGGA
ALDH4.2	NM_003748	NM_003748.2	ALDH4A1	2092	GGACAGGGTAAGA CCGTGAT	AACCGGAAGAAG TCGATGAG	CTGCAGCGTCAATCT CCGCTTG
ALDH6A1.1	NM_005589	NM_005589.2	ALDH6A1	6114	GGCTCTTTCAACAG CAGTCC	GCATGCTCCACC AGCTCT	CAGCCACTTCTTGGC TTCTCCAC
ALDOA.1	NM_000034	NM_000034.2	ALDOA	3810	GCCTGTACGTGCC AGCTC	TCATCGGAGCTT GATCTCG	TGCCAGAGCCTCAA CTGTCTCTGC
ALDOB.1	NM_000035	NM_000035.2	ALDOB	6321	CCCTCTACCAGAAG GACAGC	TAAGTTGATTCCC ACCACGA	TCCCCTTTTCTTGA GGATGTTTCTG
ALOX12.1	NM_000697	NM_000697.1	ALOX12	3861	AGTTCCTCAATGGT GCCAAC	AGCACTAGCCTG GAGGGC	CATGCTGTTGAGAC GCTCGACCTC
ALOX5.1	NM_000698	NM_000698.2	ALOX5	4259	GAGCTGCAGGACT TCGTGA	GAAGCCTGAGGA CTTGCG	CCGCATGCCGTACA CGTAGACATC
AMACR1.1	NM_014324	NM_014324.4	AMACR	3930	GGACAGTCAGTTTT AGGGTTGC	GACAGCCCAGAG ACCCAC	CAGTAACTCGGGGC CTGTTTCCC
ANGPT1.1	NM_001146	NM_001146.3	ANGPT1	2654	TCTACTTGGGGTGA CAGTGC	CCTTTTTAAAGCC CGACAGT	TCACGTGGCTCGAC TATAGAAAACCCA

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
ANGPT2.1	NM_001147	NM_001147.1	ANGPT2	2655	CCGTGAAAGCTGCTCTGTAA	TTGCAGTGGGAA GAACAGTC	AAGCTGACACAGCC CTCCCAAGTG
ANGPTL2.1	NM_012098	NM_012098.2	ANGPTL2	3982	GCCATCTGCGTCAACTCC	TAGCTCCTGCTTAGTCACTCG	TCTCCAGAAGCACCT CAGGCTCCT
ANGPTL3.3	NM_014495	NM_014495.2	ANGPTL3	6505	GTTGCGATTACTGGCAATGT	TGCTTTGTGATCCCAAGTAGA	CCAATGCAATCCCG GAAAACAAAG
ANGPTL4.1	NM_016109	NM_016109.2		3237	ATGACCTCAGATGGAGGCTG	CCGGTTGAAGTCACTGAG	CATCGTGGCGCCTC TGAATTACTG
ANGPTL7.1	NM_021146	NM_021146.2	ANGPTL7	6115	CTGCACAGACTCCAACCTCA	GCCATCCAGGTGCTTATTGT	TCACCCAGGCGGTA GTACACTCCA
ANTXR1.1	NM_032208	NM_032208.1	ANTXR1	3363	CTCCAGGTGTACCTCCAACC	GAGAAGGCTGGGAGACTCTG	AGCCTTCTCCACAG CTGCCTACA
ANXA1.2	NM_000700	NM_000700.1	ANXA1	1907	GCCCCATCCTACCTTCAATCC	CCTTTAACCATTATGGCCTTATGC	TCCTCGGATGTCGCT GCCT
ANXA2.2	NM_004039	NM_004039.1	ANXA2	2269	CAAGACACTAAGGCGACTACCA	CGTGTGCGGGCTTCAGTCAT	CCACCACACAGGTA CAGCAGCGCT
ANXA4.1	NM_001153	NM_001153.2	ANXA4	3984	TGGGAGGGATGAAGGAAAT	CTCATACAGGTCTGGGCA	TGTCTCACGAGAGC ATCGTCCAGA
ANXA5.1	NM_001154	NM_001154.2	ANXA5	3785	GCTCAAGCCTGGAGATGAC	AGAACCACCAACATCCGCT	AGTACCCTGAAGTGT CCCCCACCA
AP-1 (JUN official).2	NM_002228	NM_002228.2	JUN	2157	GACTGCAAAGATGGAAACGA	TAGCCATAAGGTCGCTCTC	CTATGACGATGCCCT CAACGCCTC
AP1M2.1	NM_005498	NM_005498.3	AP1M2	5104	ACAACGACCGCACATCT	CTGAGGCGGTATGACATGAG	CTTCATCCCGCCTGA TGGTGACTT
APAF1.2	NM_181861	NM_181861.1	APAF1	4086	CACAAGGAAGAAGCTGGTGA	CATCCTGGTTCACCTTTCAA	TGCAATTGAGCAGAA GCTCTCCAAA
APC.4	NM_000038	NM_000038.1	APC	41	GGACAGCAGGAATGTGTTTC	ACCCACTCGATTTGTTTCTG	CATTGGCTCCCCGT GACCTGTA
APOC1.3	NM_001645	NM_001645.3	APOC1	6608	CCAGCCTGATAAAGGTCCTG	CACTCTGAATCCTTGCTGGA	AGGACAGGACCTCC CAACCAAGC
APOE.1	NM_000041	NM_000041.2	APOE	4340	GCCTCAAGAGCTGGTTCG	CCTGCACCTTCTCACCA	ACTGGCGCTGCATG TCTTCCAC
APOL1.1	NM_003661	NM_003661.2	APOL1	6117	CGGACCAAGAAGTGTGACC	ATTTTGTCTGCTGGCCCTG	AGGCATATCTCTCCT GGTGGCTGC
APOLD1.1	NM_030817	NM_030817.1	APOLD1	6118	GAGCAGCTGGAGTCTCGG	AGAGATCTTGAGGTCGTGGC	CAGCTCTGCACCAA GTCCAGTCGT

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
AQP1.1	NM_198098	NM_198098.1	AQP1	5294	GCTTGCTGTATGAC CCCTG	AAGGCTGACCTCT CCCCTC	ACAGCCTTCCCTCTG CATTGACCT
AREG.2	NM_001657	NM_001657.1	AREG	87	TGTGAGTGAAATGC CTTCTAGTAGTGA	TTGTGGTTCGTTA TCATACTCTTCTG	CCGTCCTCGGGAGC CGACTATGA
ARF1.1	NM_001658	NM_001658.2	ARF1	2776	CAGTAGAGATCCCC GCAACT	ACAAGCACATGG CTATGGAA	CTTGTCTTGGGTCA CCCTGCA
ARG99.1	NM_031920	NM_031920.2		3873	GCATGGGCTACTG CATCC	CCACATCGATTCA GCCAAG	AGCTTGCTCAGTCC GTGCACAAAA
ARGHEF18.1	NM_015318	NM_015318.2	ARHGEF18	3008	ACTCTGCTTCCCAA GGGC	GAAGCTAGAGGC CCGCTC	CTGTTACACGCTCA GCCTGTCTG
ARHA.1	NM_001664	NM_001664.1	RHOA	2981	GGTCCTCCGTCGG TTCTC	GTCGCAAACCTCG GAGACG	CCACGGTCTGGTCTT CAGCTACCC
ARHGDIB.1	NM_001175	NM_001175.4	ARHGDIB	3987	TGGTCCCTAGAACA AGAGGC	TGATGGAGGATC AGAGGGAG	TAAACCGGGGCTTTC ACCCAACCT
ARRB1.1	NM_004041	NM_004041.2	ARRB1	2656	TGCAGGAACGCCT CATCAA	GGTTTGGAGGGA TCTCAAAGG	CTGGGCGAGCACGC TTACCCTTTC
ASS1.1	NM_054012	NM_054012.3	ASS1	6328	CCCCAGATAAAGG TCATTG	TGCGTACTCCATC AGGTCAT	TCTACAACCGGTTCA AGGGCCG
ATP1A1.1	NM_000701	NM_000701.6	ATP1A1	6119	AGAACGCCTATTTG GAGCTG	GGCAGAAAGAGG TGGCAG	ACCTAGGACTCGTTC TCCGAGGCC
ATP5E.1	NM_006886	NM_006886.2	ATP5E	3535	CCGCTTTCGCTACA GCAT	TGGGAGTATCGG ATGTAGCTG	TCCAGCCTGTCTCCA GTAGGCCAC
ATP6V1B1.1	NM_001692	NM_001692.3	ATP6V1B1	6293	AACCATGGGGAAC GTCTG	GGTGATGATCCG CTCGAT	CTTCCTGAACTTGGC CAATGACCC
AXL.1	NM_001699	NM_001699.3	AXL	3989	TTGCAGCCCTGTCT TCCTAC	CTGCACAGAGAA GGGGAGG	TATCCCACCTCCATC CCAGACAGG
AZU1.1	NM_001700	NM_001700.3	AZU1	6120	CCGAGGCCCTGAC TTCTT	GTCCCGGGTTGT TGAGAA	CCATCGATCCAGTCT CGGAAGAGC
B-Catenin.3	NM_001904	NM_001904.1	CTNNB1	769	GGCTCTTGTCGTA CTGTCCTT	TCAGATGACGAA GAGCACAGATG	AGGCTCAGTGATGT CTTCCCTGTCACCAG
B2M.4	NM_004048	NM_004048.1	B2M	467	GGGATCGAGACAT GTAAGCA	TGGAATTCATCCA ATCCAAAT	CGGCATCTTCAAACC TCCATGATG
BAD.1	NM_032989	NM_032989.2	BAD	7209	GGGTCAGGGGCCT CGAGAT	CTGCTCACTCGG CTCAAACCTC	TGGGCCCAGAGCAT GTTCCAGATC
BAG1.2	NM_004323	NM_004323.2	BAG1	478	CGTTGTCAGCACTT GGAATACAA	GTTCAACCTCTTC CTGTGGACTGT	CCCAATTAACATGAC CCGGCAACCAT

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
BAG2.1	NM_004282	NM_004282.2	BAG2	2808	CTAGGGGGCAAAAA GCATGA	CTAAATGCCCAAG GTGACTG	TTCCATGCCAGACAG GAAAAAGCA
Bak.2	NM_001188	NM_001188.1	BAK1	82	CCATTCCCACCATT CTACCT	GGGAACATAGAC CCACCAAT	ACACCCAGACGTC CTGGCCT
Bax.1	NM_004324	NM_004324.1	BAX	10	CCGCCGTGGACAC AGACT	TTGCCGTCAGAAA ACATGTCA	TGCCACTCGGAAAAA GACCTCTCGG
BBC3.2	NM_014417	NM_014417.1	BBC3	574	CCTGGAGGGTCCT GTACAAT	CTAATTGGGCTCC ATCTCG	CATCATGGGACTCCT GCCCTTACC
Bcl2.2	NM_000633	NM_000633.1	BCL2	61	CAGATGGACCTAGT ACCCACTGAGA	CCTATGATTTAAG GGCATTITTC	TTCCACGCCGAAGG ACAGCGAT
BCL2A1.1	NM_004049	NM_004049.2	BCL2A1	6322	CCAGCCTCCATGTA TCATCA	TGAAGCTGTTGA GGCAATGT	CAGTCAAGCTCAGT GAGCATTCTCAGC
BCL2L12.1	NM_138639	NM_138639.1	BCL2L12	3364	AACCCACCCCTGTC TTGG	CTCAGCTGACGG GAAAGG	TCCGGGTAGCTCTC AAACTCGAGG
Bclx.2	NM_001191	NM_001191.1	BCL2L1	83	CTTTTGTGGAAGTC TATGGGAACA	CAGCGGTTGAAG CGTTCCT	TTCGGCTCTCGGCT GCTGCA
BCRP.1	NM_004827	NM_004827.1	ABCG2	364	TGTAAGTGGGAAGA ATATTTGGTAAA	GCCACGTGATTCT TCCACAA	CAGGGCATCGATCT CTCACCTGG
BFGF.3	NM_007083	NM_007083.1	NUDT6	345	CCAGGAAGAATGCT TAAGATGTGA	TGGTGATGGGAG TTGTATTTTCAG	TTGCCAGGTCATTG AGATCCATCCA
BGN.1	NM_001711	NM_001711.3	BGN	3391	GAGCTCCGCAAGG ATGAC	CTTGTTGTTCCACC AGGACGA	CAAGGGTCTCCAGC ACCTCTACGC
BHLHB3.1	NM_030762	NM_030762.1	BHLHE41	6121	AGGAAGATCCCTC GCAGC	TTGAACCTCCGTC CTTCG	AGGAAGCTCCCTGA ATCCTTGCGT
BIK.1	NM_001197	NM_001197.3	BIK	2281	ATTCCTATGGCTCT GCAATTGTC	GGCAGGAGTGAA TGGCTCTTC	CCGGTTAACTGTGG CCTGTGCC
BIN1.3	NM_004305	NM_004305.1	BIN1	941	CCTGCAAAAGGGAA CAAGAG	CGTGGTTGACTCT GATCTCG	CTTCGCCTCCAGATG GCTCCC
BLR1.1	NM_001716	NM_001716.3	CXCR5	6280	GACCAAGCAGGAA GCTCAGA	AGCGCTGTTTCG GTCAGA	CCAGGGGCAGCTAC CTGAACTCAA
BNIP3.1	NM_004052	NM_004052.2	BNIP3	3937	CTGGACGGAGTAG CTCCAAG	GGTATCTTGTTGGT GTCTGCG	CTCTCACTGTGACAG CCCACCTCG
BRCA1.1	NM_007294	NM_007294.3	BRCA1	7481	TCAGGGGGGCTAGA AATCTGT	CCATTCCAGTTGA TCTGTGG	CTATGGGCCCTTCAC CAACATGC
BTRC.1	NM_033637	NM_033637.2	BTRC	2555	GTTGGGACACAGTT GGTCTG	TGAAGCAGTCAGT TGTGCTG	CAGTCGGCCCCAGGA CGGTCTACT

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
BUB1.1	NM_004336	NM_004336.1	BUB1	1647	CCGAGGTTAATCCA GCACGTA	AAGACATGGCGC TCTCAGTTC	TGCTGGGAGCCTAC ACTTGGCCC
BUB3.1	NM_004725	NM_004725.1	BUB3	3016	CTGAAGCAGATGGT TCATCATT	GCTGATTCCCAAG AGTCTAACC	CCTCGCTTTGTTTAA CAGCCCAGG
c-kit.2	NM_000222	NM_000222.1	KIT	50	GAGGCAACTGCTTA TGGCTTAATTA	GGCACTCGGCTT GAGCAT	TTACAGCGACAGTCA TGGCCGCAT
C13orf15.1	NM_014059	NM_014059.2	C13orf15	6122	TAGAATCTGCTGCC AGAGGG	CAAGGGCTGATTT TAAGGTGA	TGCACTCAACCTTCT ACCAGGCCA
C1QA.1	NM_015991	NM_015991.2	C1QA	6123	CGGTCATCACCAAC CAGG	CGGGTACAGTGC AGACGA	AGAACCGTACCAGAA CCACTCCGG
C1QB.1	NM_000491	NM_000491.3	C1QB	6124	CCAGTGGCCTCAC AGGAC	CCCATGGGATCTT CATCATC	TCCAGGAGGCGTC TGACACAGTA
C20 orf1.1	NM_012112	NM_012112.2	TPX2	1239	TCAGCTGTGAGCTG CGGATA	ACGGTCCTAGGT TTGAGGTAAAGA	CAGGTCCCATTGCC GGGCG
C3.1	NM_000064	NM_000064.2	C3	6125	CGTGAAGGAGTGC AGAAAGA	ACTCGGTGTCCG GGAATT	ACATCCCACCTGCAG ACCTCAGTG
C3AR1.1	NM_004054	NM_004054.2	C3AR1	6126	AAGCCGCATCCCA GACTT	TGTTAAGTGCCCT TGCTGG	CAACCCCCAGAGATT CCGATTCAG
C7.1	NM_000587	NM_000587.2	C7	6127	ATGTCTGAGTGTGA GGCGG	AGGCCTTATGCT GGTGACAG	ATGCTCTGCCCTCTG CATCTCAGA
CA12.1	NM_001218	NM_001218.3	CA12	6128	CTCTCTGAAGGTGT CCTGGC	ACAGGAACTGAG GGGTGCT	AGACACCAGTGCTTC TCCAGGGCT
CA2.1	NM_000067	NM_000067.1	CA2	5189	CAACGTGGAGTTTG ATGACTCT	CTGTAAGTGCCAT CCAGGG	CCTCCCTTGAGCACT GCTTTGTCC
CA9.3	NM_001216	NM_001216.1	CA9	482	ATCCTAGCCCTGGT TTTTGG	CTGCCTTCTCATC TGCACAA	TTTGCTGTACCAGC GTCGC
CACNA2D1.1	NM_000722	NM_000722.2	CACNA2D1	6129	CAAACATTAGCTGG GCCTGT	CAGCCAGTGGGT GCCTTA	CCATGGCATAACACT AAGGCGCAG
CALD1.2	NM_004342	NM_004342.4	CALD1	1795	CACTAAGGTTTGAG ACAGTTCAGAA	GCGAATTAGCCCT CTACAAGTGA	AACCCAAGCTCAAGA CGCAGGACGAG
CASP1.1	NM_033292	NM_033292.2	CASP1	6132	AGAAAGCCACATA GAGAAGGA	TGTGGGATGTCT CCAAGAAA	CGCTTTCTGCTCTTC CACACCAGA
CASP10.1	NM_001230	NM_001230.4	CASP10	6133	ACCTTTCTCTTGGC CGGAT	GTGGGGACTGTC CACTGC	TCTACTGCATCTGCC AGCCCTGAG
CASP6.1	NM_032992	NM_032992.2	CASP6	6134	CCTCACACTGGTGA ACAGGA	AATTGCACTTGGG TCTTTGC	AAAGTCCACTCGGC GCTGAGAAAC

Table A

Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
Caspase 3.1	NM_032991	NM_032991.2	CASP3	5963	TGAGCCTGAGCAG AGACATGA	CCTTCCTGCGTG GTCCAT	TCAGCCTGTTCCATG AAGGCAGAGC
CAT.1	NM_001752	NM_001752.1	CAT	2745	ATCCATTTCGATCTC ACCAAGGT	TCCGGTTTAAAGAC CAGTTTACCA	TGGCCTCACAAGGA CTACCCTCTCATCC
CAV1.1	NM_001753	NM_001753.3	CAV1	2557	GTGGCTCAACATTG TGTTCC	CAATGGCCTCCAT TTTACAG	ATTTCAGCTGATCAG TGGGCCTCC
CAV2.1	NM_198212	NM_198212.1	CAV2	6460	CTTCCTGGGACG ACTTG	CTCCTGGTCACC CTTCTGG	CCCGTACTGTCATGC CTCAGAGCT
CCL18.1	NM_002988	NM_002988.2	CCL18	3994	GCTCCTGTGCACAA GTTGG	TGGAATCTGCCA GGAGGTA	CAACAAAGAGCTCTG CTGCCTCGT
CCL19.1	NM_006274	NM_006274.2	CCL19	4107	GAACGCATCATCCA GAGACTG	CCTCTGCACGGT CATAGGTT	CGCTTCATCTTGGCT GAGGTCCTC
CCL20.1	NM_004591	NM_004591.1	CCL20	1998	CCATGTGCTGTACC AAGAGTTTG	CGCCGCAGAGGT GGAGTA	CAGCACTGACATCAA AGCAGCCAGGA
CCL4.2	NM_002984	NM_002984.1	CCL4	4148	GGGTCCAGGAGTA CGTGTATGAC	CCTTCCCTGAAGA CTTCCTGTCT	ACTGAACTGAGCTG CTCA
CCL5.2	NM_002985	NM_002985.2	CCL5	4088	AGGTTCTGAGCTCT GGCTTT	ATGCTGACTTCCT TCCTGGT	ACAGAGCCCTGGCA AAGCCAAG
CCNB1.2	NM_031966	NM_031966.1	CCNB1	619	TTCAGGTTGTTGCA GGAGAC	CATCTTCTTGGGC ACACAAT	TGTCTCCATTATTGA TCGGTTCATGCA
CCND1.3	NM_053056	NM_001758.1	CCND1	88	GCATGTTTCGTGGC CTCTAAGA	CGGTGTAGATGC ACAGCTTCTC	AAGGAGACCATCCC CCTGACGGC
CCNE1.1	NM_001238	NM_001238.1	CCNE1	498	AAAGAAGATGATGA CCGGGTTTAC	GAGCCTCTGGAT GGTGCAAT	CAAACCTAACGTGCA AGCCTCGGA
CCNE2 variant 1.1	NM_057749	NM_057749var1	CCNE2	1650	GGTCACCAAGAAAC ATCAGTATGAA	TTCAATGATAATG CAAGGACTGATC	CCCAGATAATACAGG TGGCCAACAATTCCT
CCNE2.2	NM_057749	NM_057749.1	CCNE2	502	ATGCTGTGGCTCCT TCCTAACT	ACCCAAATTGTGA TATACAAAAAGGT T	TACCAAGCAACCTAC ATGTCAAGAAAGCCC
CCR1.1	NM_001295	NM_001295.2	CCR1	6135	TCCAAGACCCAATG GGAA	TCGTAGGCTTTTCG TGAGGA	ACTCACCACACCTGC AGCCTTCAC
CCR2.1	NM_000648	NM_000648.1		4109	CTCGGGAATCCTGA AAACC	GACTCTCACTGCC CTATGCC	TCTTCTCGTTTCGAC ACCGAAGCA
CCR4.2	NM_005508	NM_005508.4	CCR4	6502	AGACCCTGGTGGA GCTAGAA	AGAGTTTCTGTGG CCTGGAT	TCCTTCAGGACTGCA CCTTTGAAAGA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
CCR5.1	NM_000579	NM_000579.1	CCR5	4119	CAGACTGAATGGG GGTGG	CTGGTTTGTCTGG AGAAGGC	TGGAATAAGTACCTA AGGCGCCCCC
CCR7.1	NM_001838	NM_001838.2	CCR7	2661	GGATGACATGCACT CAGCTC	CCTGACATTTCCC TTGTCCT	CTCCCATCCCAGTG GAGCCAA
CD105.1	NM_000118	NM_000118.1	ENG	486	GCAGGTGTCAGCA AGTATGATCAG	TTTTTCCGCTGTG GTGATGA	CGACAGGATATTGAC CACCGCCTCATT
CD14.1	NM_000591	NM_000591.1	CD14	4341	GTGTGCTAGCGTAC TCCCG	GCATGGTGCCGG TTATCT	CAAGGAACTGACGC TCGAGGACCT
CD18.2	NM_000211	NM_000211.1	ITGB2	49	CGTCAGGACCCAC CATGTCT	GGTTAATTGGTGA CATCCTCAAGA	CGCGGCCGAGACAT GGCTTG
CD1A.1	NM_001763	NM_001763.1	CD1A	4166	GGAGTGGGAAGGAA CTGGA	TCATGGGCGTAT CTACGAAT	CGCACCATTTCGGTC ATTTGAGG
CD24.1	NM_013230	NM_013230.1	CD24	2364	TCCAATAATGCCA CCACCAA	GAGAGAGTGAGA CCACGAAGAGAC T	CTGTTGACTGCAGG GCACCACCA
CD274.2	NM_014143	NM_014143.2	CD274	4076	GCTGCATGATCAGC TATGGT	TGTTGTATGGGG CATTGACT	CACAGTAATTCGCTT GTAGTCGGCACC
CD31.3	NM_000442	NM_000442.1	PECAM1	485	TGTATTTCAAGACC TCTGTGCACTT	TTAGCCTGAGGAA TTGCTGTGTT	TTTATGAACCTGCCC TGCTCCACA
CD34.1	NM_001773	NM_001773.1	CD34	3814	CCACTGCACACACC TCAGA	CAGGAGTTTACCT GCCCT	CTGTTCTTGGGGCC CTACACCTTG
CD36.1	NM_000072	NM_000072.2	CD36	6138	GTAACCCAGGACG CTGAGG	AAGGTTCTGAAGAT GGCACC	CACAGTCTCTTTCT GCAGCCCAA
CD3z.1	NM_000734	NM_000734.1	CD247	64	AGATGAAGTGGAA GGCGCTT	TGCCTCTGTAATC GGCAACTG	CACCGCGGCCATCC TGCA
CD4.1	NM_000616	NM_000616.2	CD4	4168	GTGCTGGAGTCGG GACTAAC	TCCCTGCATTCAA GAGGC	CAGGTCCCTTGTC CAAGTTCCAC
CD44.1	NM_000610	NM_000610.3	CD44	4267	GGCACCCTGCTTA TGAAGG	GATGCTCATGGT GAATGAGG	ACTGGAACCCAGAA GCACACCCTC
CD44s.1	M59040	M59040.1		1090	GACGAAGACAGTC CCTGGAT	ACTGGGGTGGAA TGTGTCTT	CACCGACAGCACAG ACAGAATCCC
CD44v6.1	AJ251595v6	AJ251595v6		1061	CTCATACCAGCCAT CCAATG	TTGGGTTGAAGAA ATCAGTCC	CACCAAGCCCAGAG GACAGTTCCT
CD53.1	NM_000560	NM_000560.3	CD53	6139	CGACAGCATCCACC GTTAC	TGCAGAAATGACT GGATGGA	CACGCTGCCTTGGT GCTATTGTCT
CD68.2	NM_001251	NM_001251.1	CD68	84	TGGTTCCCAGCCCT GTGT	CTCCTCCACCCTG GGTTGT	CTCCAAGCCCAGATT CAGATTCGAGTCA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
CD82.3	NM_002231	NM_002231.2	CD82	316	GTGCAGGCTCAGG TGAAGTG	GACCTCAGGGCG ATTCAATGA	TCAGCTTCTACAACT GGACAGACAACGCT G
CD8A.1	NM_171827	NM_171827.1	CD8A	3804	AGGGTGAGGTGCT TGAGTCT	GGGCACAGTATC CCAGGTA	CCAACGGCAAGGGA ACAAGTACTTCT
CD99.1	NM_002414	NM_002414.3	CD99	6323	GTTCCCTCCGGTAGC TTTTCA	ACCATCACTGCCT CCTTTTC	TCCACCTGAAACGCC ATCCG
cdc25A.4	NM_001789	NM_001789.1	CDC25A	90	TCTTGCTGGCTACG CCTCTT	CTGCATTGTGGC ACAGTTCTG	TGTCCCTGTTAGACG TCCTCCGTCCATA
CDC25B.1	NM_021873	NM_021874.1	CDC25B	389	AAACGAGCAGTTTG CCATCAG	GTTGGTGATGTTT CGAAGCA	CCTCACCGGCATAG ACTGGAAGCG
CDH1.3	NM_004360	NM_004360.2	CDH1	11	TGAGTGTCCCCCG GTATCTTC	CAGCCGCTTTCA GATTTTCAT	TGCCAATCCCGATGA AATTGGAAATTT
CDH13.1	NM_001257	NM_001257.3	CDH13	6140	GCTACTTCTCCACT GTCCCCG	CCTCTCTGTGGA CCTGCCT	AGTCTGAATGCTGCC ACAACCAGC
CDH16.1	NM_004062	NM_004062.2	CDH16	4529	GACTGTCTGAATGG CCCAG	CCAGGGGACTCA GATGGA	CAGAGGCCAAGCTC CCAGCTAGAG
CDH2.1	NM_001792	NM_001792.2	CDH2	3965	TGACCGATAAGGAT CAACCC	GATCTCCGCCACT GATTCTG	ATACACCAGCCTGGA ACGCAGTGT
CDH5.1	NM_001795	NM_001795.2	CDH5	6142	ACAGGAGACGTGTT CGCC	CAGCAGTGAGGT GGTACTCTGA	TATTCTCCCGGTCCA GCCTCTCAA
CDH6.1	NM_004932	NM_004932.2	CDH6	3998	ACACAGGCGACATA CAGGC	CTCGAAGGATGTA AACGGGT	TTTCTTCCCTGTCCA GCCTCTTGG
CDK4.1	NM_000075	NM_000075.2	CDK4	4176	CCTTCCCATCAGCA CAGTTC	TTGGGATGCTCAA AAGCC	CCAGTCGCCTCAGT AAAGCCACCT
CDK6.1	NM_001259	NM_001259.5	CDK6	6143	AGTGCCCTGTCTCA CCCA	GCAGGTGGGAAT CCAGGT	TCTTTGCACCTTTCC AGGTCCTGG
CDKN2A.2	NM_000077	NM_000077.3	CDKN2A	4278	AGCACTCACGCCCT AAGC	TCATGAAGTCGAC AGCTTCC	CGCAAGAAATGCCC ACATGAATGT
CEACAM1.1	NM_001712	NM_001712.2	CEACAM1	2577	ACTTGCTGTTCAG AGCACTCA	TGGCAAATCCGAA TTAGAGTGA	TCCTTCCCACCCCA GTCCTGTC
CEBPA.1	NM_004364	NM_004364.2	CEBPA	2961	TTGGTTTTTGCTCGG ATACTTG	GTCTCAGACCCTT CCCCC	AAAATGAGACTCTCC GTCGGCAGC
CENPF.1	NM_016343	NM_016343.2	CENPF	3251	CTCCCGTCAACAGC GTTC	GGGTGAGTCTGG CCTTCA	AACTGGACCAGGA GTGCATCCAG
CFLAR.1	NM_003879	NM_003879.3	CFLAR	6144	GGACTTTTGTCCAG TGACAGC	CGGCGCTTCTCT CCTACA	CTCCTCCCGTGGTC CTTGTTGTCT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
CGA (CHGA official).3	NM_001275	NM_001275.2	CHGA	1132	CTGAAGGAGCTCC AAGACCT	CAAAACCGCTGT GTTTCTTC	TGCTGATGTGCCCTC TCCTTGG
Chk1.2	NM_001274	NM_001274.1	CHEK1	490	GATAAATTGGTACA AGGGATCAGCTT	GGGTGCCAAGTA ACTGACTATTCA	CCAGCCCACATGTC CTGATCATATGC
Chk2.3	NM_007194	NM_007194.1	CHEK2	494	ATGTGGAACCCCCA CCTACTT	CAGTCCACAGCA CGGTTATACC	AGTCCCAACAGAAAC AAGAACTTCAGGCG
CIAP1.2	NM_001166	NM_001166.2	BIRC2	326	TGCCTGTGGTGGG AAGCT	GGAAAATGCCTC CGGTGTT	TGACATAGCATCATC CTTTGGTTCCAGTT
ciAP2.2	NM_001165	NM_001165.2	BIRC3	79	GGATATTTCCGTGG CTCTTATTCA	CTTCTCATCAAGG CAGAAAAATCTT	TCTCCATCAAATCCT GTAAACTCCAGAGCA
CLCNKB.1	NM_000085	NM_000085.1	CLCNKB	4308	GTGACCCTGAAGCT GTCCC	GGTTCAACAGCTC AAAGAGGTT	AGAGACTTCCCTGCA TGAGGCACA
CLDN10.1	NM_182848	NM_182848.2	CLDN10	6145	GGTCTGTGGATGAA CTGCG	GATAGTAAAATGC GGTCGGC	TGGAAAGAACCCCAAC GCGTTACCT
CLDN7.2	NM_001307	NM_001307.3	CLDN7	2287	GGTCTGCCCTAGTC ATCCTG	GTACCCAGCCTT GCTCTCAT	TGCACTGCTCTCCTG TTCCTGTCC
CLU.3	NM_001831	NM_001831.1	CLU	2047	CCCCAGGATACCTA CCACTACCT	TGCGGGACTTGG GAAAGA	CCCTTCAGCCTGCC CCACCG
cMet.2	NM_000245	NM_000245.1	MET	52	GACATTTCCAGTCC TGCAGTCA	CTCCGATCGCAC ACATTTGT	TGCCTCTGTGCCCCA CCCTTTGT
cMYC.3	NM_002467	NM_002467.1	MYC	45	TCCCTCCACTCGGA AGGACTA	CGGTTGTTGCTG ATCTGTCTCA	TCTGACACTGTCCAA CTTGACCCTCTT
COL18A1.1	NM_030582	NM_030582.3	COL18A1	6146	AGCTGCCATCACG CCTAC	GTGGCTACTTGG AGGCAGTC	CGTGCTCTGCATTGA GAACAGCTTC
COL1A1.1	NM_000088	NM_000088.2	COL1A1	1726	GTGGCCATCCAGC TGACC	CAGTGGTAGGTG ATGTTCTGGGA	TCCTGCGCCTGATGT CCACCG
COL1A2.1	NM_000089	NM_000089.2	COL1A2	1727	CAGCCAAGAACTG GTATAGGAGCT	AAACTGGCTGCC AGCATTG	TCTCCTAGCCAGAC GTGTTTCTTGTCTT G
COL4A1.1	NM_001845	NM_001845.4	COL4A1	6147	ACAAAGGCCTCCCA GGAT	GAGTCCCAGGAA GACCTGCT	CTCCTTTGACACCAG GGATGCCAT
COL4A2.1	NM_001846	NM_001846.2	COL4A2	6148	CAACCCTGGTGATG TCTGC	CGCAGTGGTAGA GAGCCAGT	ACTATGCCAGCCGG AACGACAAGT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
COL5A2.2	NM_000393	NM_000393.3	COL5A2	6653	GGTCGAGGAACCC AAGGT	GCCTGGAGGTCC AACTCTG	CCAGGAAATCCTGTA GCACCAGGC
COL7A1.1	NM_000094	NM_000094.2	COL7A1	4984	GGTGACAAAGGAC CTCGG	ACCAGGCTCTCC CTTGCT	CTTGTCACCAGGGT CCCCATTGTC
COX2.1	NM_000963	NM_000963.1	PTGS2	71	TCTGCAGAGTTGGA AGCACTCTA	GCCGAGGCTTTT CTACCAGAA	CAGGATACAGCTCCA CAGCATCGATGTC
CP.1	NM_000096	NM_000096.1	CP	3244	CGTGAGTACACAGA TGCCTCC	CCAGGATGCCAA GATGCT	TCTTCAGGGCCTCTC TCCTTTCGA
CPB2.1	NM_001872	NM_001872.3	CPB2	6294	GGCACATACGGATT CTTGCT	CAGCGGCAAAAG CTTCTCTA	CGGAGCGTTACATC AAACCCACCT
CRADD.1	NM_003805	NM_003805.3	CRADD	6149	GATGGTGCCTCCA GCAAC	GAGTGAAAGTCA GGATTCAGCC	CATGACTCAGGGAC ACACTCCCCA
cripto (TDGF1 official).1	NM_003212	NM_003212.1	TDGF1	1096	GGGTCTGTGCCCC ATGAC	TGACCGTGCCAG CATTTACA	CCTGGCTGCCCAAG AAGTGTTCCCT
CRP.1	NM_000567	NM_000567.2	CRP	4187	GACGTGAACCACA GGGTGT	CTCCAGATAGGG AGCTGGG	CTGTCAGAGGAGCC CATCTCCCAT
CSF1.1	NM_000757	NM_000757.3	CSF1	510	TGCAGCGGCTGATT GACA	CAACTGTTCTCTGG TCTACAAACTCA	TCAGATGGAGACCT CGTGCCAAATTACA
CSF1R.2	NM_005211	NM_005211.1	CSF1R	2289	GAGCACAACCAAAC CTACGA	CCTGCAGAGATG GGTATGAA	AGCCACTCCCCACG CTGTTGT
CSF2.1	NM_000758	NM_000758.2	CSF2	2779	GAACCTGAAGGACT TTCTGCTTGT	CTCATCTGGCCG GTCTCACT	ATCCCCTTTGACTGC TGGGAGCCAG
CSF2RA.2	NM_006140	NM_006140.3	CSF2RA	4477	TACCACACCCAGCA TTCCTC	CTAGAGGCTGGT GCCACTGT	CGCAGATCCGATTTC TCTGGGATC
CSF3.2	NM_000759	NM_000759.1	CSF3	377	CCCAGGCCTCTGT GTCCTT	GGAGGACAGGAG CTTTTTCTCA	TGCATTTCTGAGTTT CATTCTCCTGCCTG
CTGF.1	NM_001901	NM_001901.1	CTGF	2153	GAGTTCAAGTGCCC TGACG	AGTTGTAATGGCA GGCACAG	AACATCATGTTCTTC TTCATGACCTCGC
CTSB.1	NM_001908	NM_001908.1	CTSB	382	GGCCGAGATCTAC AAAAACG	GCAGGAAGTCCG AATACACA	CCCCGTGAGGGAG CTTTCTC
CTSD.2	NM_001909	NM_001909.1	CTSD	385	GTACATGATCCCCT GTGAGAAGGT	GGGACAGCTTGT AGCCTTTGC	ACCCTGCCCGOGAT CACTACTGA
CTSH.2	NM_004390	NM_004390.1	CTSH	845	GCAAGTTCCAACCT GGAAAG	CATCGCTTCCTCG TCATAGA	TGGCTACATCCTTGA CAAAGCCGA
CTSL.2	NM_001912	NM_001912.1	CTSL1	446	GGGAGGCTTATCTC ACTGAGTGA	CCATTGCAGCCTT CATTGC	TTGAGGCCAGAGC AGTCTACCAGATTCT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
CTSL2.1	NM_001333	NM_001333.2	CTSL2	1667	TGTCTCACTGAGCG AGCAGAA	ACCATTGCAGCC CTGATTG	CTTGAGGACGCGAA CAGTCCACCA
CTSS.1	NM_004079	NM_004079.3	CTSS	1799	TGACAACGGCTTTC CAGTACAT	TCCATGGCTTTGT AGGGATAGG	TGATAACAAGGGCAT CGACTCAGACGCT
CUBN.1	NM_001081	NM_001081.2	CUBN	4110	GAGGCCGTTACTGT GGCA	GAATCTCAGCGTC AGGGC	TGCCCCATCCTATCA CATCCTTCA
CUL1.1	NM_003592	NM_003592.2	CUL1	1889	ATGCCCTGGTAATG TCTGCAT	GCGACCACAAGC CTTATCAAG	CAGCCACAAAGCCA GCGTCATTGT
CUL4A.1	NM_003589	NM_003589.1	CUL4A	2780	AAGCATCTTCCTGT TCTTGGA	AATCCCATATCCC AGATGGA	TATGTGCTGCAGAAC TCCACGCTG
CX3CL1.1	NM_002996	NM_002996.3	CX3CL1	6150	GACCCTTGCCGTCT ACCTG	GGAGTGTTCTTA GCACCTGG	TAGAACCCAGCCCAT AAGAGGCC
CX3CR1.1	NM_001337	NM_001337.3	CX3CR1	4507	TTCCCAGTTGTGAC ATGAGG	GCTAAATGCAACC GTCTCAGT	ACTGAGGGCCAGCC TCAGATCCT
CXCL10.1	NM_001565	NM_001565.1	CXCL10	2733	GGAGCAAAATCGAT GCAGT	TAGGGAAGTGAT GGGAGAGG	TCTGTGTGGTCCATC CTTGGAAGC
CXCL12.1	NM_000609	NM_000609.3	CXCL12	2949	GAGCTACAGATGC CCATGC	TTTGAGATGCTTG ACGTTGG	TTCTTCGAAAGCCAT GTTGCCAGA
CXCL14.1	NM_004887	NM_004887.3	CXCL14	3247	TGCGCCCTTTCTCTC TGTA	CAATGCGGCATAT ACTGGG	TACCCTTAAGAACGC CCCCTCCAC
CXCL9.1	NM_002416	NM_002416.1	CXCL9	5191	ACCAGACCATTGTC TCAGAGC	GTTACCAGAGGC TAGCCAACA	TGCTGGCTCTTTCTC GGCTACTCC
CXCR4.3	NM_003467	NM_003467.1	CXCR4	2139	TGACCGCTTCTACC CCAATG	AGGATAAGGCCA ACCATGATGT	CTGAAACTGGAACAC AACCACCCACAAG
CXCR6.1	NM_006564	NM_006564.1	CXCR6	4002	CAGAGCCTGACGG ATGTGT	GCAGAGTGCAGA CAAACACC	CTGGTGAACCTACC CCTGGCTGAC
CYP2C8.2	NM_000770	NM_000770.2	CYP2C8	505	CCGTGTTCAAGAG GAAGCTC	AGTGGGATCACA GGGTGAAG	TTTTCTCAACTCCTC CACAAGGCA
CYP2C8v2.1	NM_030878	NM_030878.1		5423	GCTGTAGTGCACG AGATCCA	CAGTGGTCACTG CATGGG	ATACAGTGACCTTGT CCCCACCGG
CYP3A4.2	NM_017460	NM_017460.3	CYP3A4	586	AGAACAAGGACAAC ATAGATCCTTACAT AT	GCAAACCTCATGC CAATGC	CACACCCTTTGGAAG TGGACCCAGAA
CYR61.1	NM_001554	NM_001554.3	CYR61	2752	TGCTCATTCTTGAG GAGCAT	GTGGCTGCATTA GTGTCCAT	CAGCACCCTTGGA GTTTCGAAAT
DAG1.1	NM_004393	NM_004393.2	DAG1	5968	GTGACTGGGCTCAT GCCT	ATCCCACTTGTGC TCCTGTC	CAAGTCAGAGTTTCC CTGGTGCC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
DAPK1.3	NM_004938	NM_004938.1	DAPK1	636	CGCTGACATCATGA ATGTTCT	TCTCTTTCAGCAA CGATGTGTCTT	TCATATCCAACTCG CCTCCAGCCG
DCBLD2.1	NM_080927	NM_080927.3	DCBLD2	3821	TCACCAGGGCAGG AAGTTTA	GGTTGCATACTCA GGCCC	CATGCCTATGCTGAA CCACTCCCA
DCC.3	NM_005215	NM_005215.1	DCC	3763	AAATGTCCTCCTCG ACTGCT	TGAATGCCATCTT TCTTCCA	ATCACTGGAACCTCT CGGTCCGAC
DCN.1	NM_001920	NM_001920.3	DCN	6151	GAAGGCCACTATCA TCCTCCT	GCCTCTCTGTTGA AACGGTC	CTGCTTGCACAAGTT TCCTGGGCT
DCXR.1	NM_016286	NM_016286.2	DCXR	6311	CCATAGCGTCTACT GCTCCA	AGCTCTAGGGCC ATCACCT	TCAGCATGTCCAGG GCACCC
DDC.1	NM_000790	NM_000790.3	DDC	5411	CAGAGCCCAGACA CCATGA	CCACGTAATCCAC CATCTCC	CCTCTCCTTCGGAAT TCACTTGCG
DEFB1.1	NM_005218	NM_005218.3	DEFB1	4124	GATGGCCTCAGGT GGTAACT	TGCTGACGCAATT GTAATGAT	CTCACAGGCCTTGG CCACAGATCT
DET1.1	NM_017996	NM_017996.2	DET1	2643	CTTGTGGAGATCAC CCAATCAG	CCCGCCTGGATC TCAAACCT	CTATGCCCGGGACT CGGGCCT
DHPS.3	NM_013407	NM_013407.1	DHPS	1722	GGGAGAACGGGAT CAATAGGAT	GCATCAGCCAGT CCTCAAACCT	CTCATTGGGCACCA GCAGGTTTCC
DIABLO.1	NM_019887	NM_019887.1	DIABLO	348	CACAATGGCGGCT CTGAAG	ACACAAACACTGT CTGTACCTGAAGA	AAGTTACGCTGCGC GACAGCCAA
DIAPH1.1	NM_005219	NM_005219.2	DIAPH1	2705	CAAGCAGTCAAGGA GAACCA	AGTTTTGCTCGCC TCATCTT	TTCTTCTGTCTCCCG CCGCTTC
DICER1.2	NM_177438	NM_177438.1	DICER1	1898	TCCAATTCCAGCAT CACTGT	GGCAGTGAAGGC GATAAAGT	AGAAAAGCTGTTTGT CTCCCCAGCA
DKFZP564O0823.1	NM_015393	NM_015393.2	PARM1	3874	CAGCTACACTGTCTG CAGTCC	ATGAGGCTGGAG CTTGAGG	TGCTGAGCCTCCCA CACTCATCTC
DLC1.1	NM_006094	NM_006094.3	DLC1	3018	GATTCAGACGAGG ATGAGCC	CACCTCTTGCTGT CCCTTTG	AAAGTCCATTGCCA CTGATGGCA
DLL4.1	NM_019074	NM_019074.2	DLL4	5273	CACGGAGGTATAA GGCAGGAG	AGAAGGAAGGTC CAGCCG	CTACCTGGACATCCC TGCTCAGCC
DPEP1.1	NM_004413	NM_004413.2	DPEP1	6295	GGACTCCAGATGC CAGGA	TAAGCCCAGGCG TCCTCT	CACATGCAAGGACC AGCATCTCCT
DPYS.1	NM_001385	NM_001385.1	DPYS	6152	AAAGAATGGCACCA TGCAG	AGTCGGGTGTTG AGGGGT	CACCATGTCATGGGT CCACCTTTG

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
DR4.2	NM_003844	NM_003844.1	TNFRSF10A	896	TGCACAGAGGGTG TGGGTTAC	TCTTCATCTGATT TACAAGCTGTACA TG	CAATGCTTCCAACAA TTTGTGTTGCTTGCC
DR5.2	NM_003842	NM_003842.2	TNFRSF10B	902	CTCTGAGACAGTGC TTCGATGACT	CCATGAGGCCCA ACTTCCT	CAGACTTGGTGCCC TTTGACTCC
DUSP1.1	NM_004417	NM_004417.2	DUSP1	2662	AGACATCAGCTCCT GGTTCA	GACAAACACCCTT CCTCCAG	CGAGGCCATTGACTT CATAGACTCCA
DUSP9.1	NM_001395	NM_001395.1	DUSP9	6324	CGTCCTAATCAACG TGCCTA	CCCGCAAAGAAAA AGTAACAG	CGCTCGGAGCCTGC CTCTTC
E2F1.3	NM_005225	NM_005225.1	E2F1	1077	ACTCCCTCTACCCT TGAGCA	CAGGCCTCAGTT CCTTCAGT	CAGAAGAACAGCTCA GGGACCCCT
EBAG9.1	NM_004215	NM_004215.3	EBAG9	4151	CGCTCCTGTTTTTC TCATCTGT	ACCGAACTGGG TGATGG	CAGTGGGTTTTGATT CCCACCATG
ECRG4.1	NM_032411	NM_032411.1	C2orf40	3869	GCTCCTGCTCCTGT GCTG	TTTTGAAGCATCA GCTTGAGTT	ATTTCACCTTATGCC ACCTGGGCC
EDG2.1	NM_001401	NM_001401.3	LPAR1	4673	ACGAGTCCATTGCC TTCTTT	GCTTGCTGACTGT GTTCCAT	CGAAGTGGAAGCA TCTTGCCACA
EDN1 endothelin.1	NM_001955	NM_001955.1	EDN1	331	TGCCACCTGGACAT CATTTG	TGGACCTAGGGC TTCCAAGTC	CACTCCCGAGCACG TTGTTCCGT
EDN2.1	NM_001956	NM_001956.2	EDN2	2646	CGACAAGGAGTGC GTCTACTTCT	CAGGCCGTAAGG AGCTGTCT	CCACTTGGACATCAT CTGGGTGAACACTC
EDNRA.2	NM_001957	NM_001957.1	EDNRA	3662	TTTCCTCAAATTTG CCTCAAG	TTACACATCCAAC CAGTGCC	CCTTGCCTCAGGG CATCCTTTT
EDNRB.1	NM_000115	NM_000115.1	EDNRB	3185	ACTGTGAACTGCCT GGTGC	ACCACAGCATGG GTGAGAG	TGCTACCTGCCCTT TGTCATGTG
EEF1A1.1	NM_001402	NM_001402.5	EEF1A1	5522	CGAGTGGAGACTG GTGTTCTC	CCGTTGTAACGTT GACTGGA	CAAAGGTGACCACC ATACCGGGTT
EFNB1.2	NM_004429	NM_004429.3	EFNB1	3299	GGAGCCCGTATCC TGGAG	GGATAGATCACCA AGCCCTTC	CCCTCAACCCCAAGT TCCTGAGTG
EFNB2.1	NM_004093	NM_004093.2	EFNB2	2597	TGACATTATCATCC CGCTAAGGA	GTAGTCCCCGCT GACCTTCTC	CGGACAGCGTCTTC TGCCCTCACT
EGF.3	NM_001963	NM_001963.2	EGF	158	CTTTGCCTTGCTCT GTCACAGT	AAATACCTGACAC CCTTATGACAAAT T	AGAGTTTAACAGCCC TGCTCTGGCTGACTT
EGFR.2	NM_005228	NM_005228.1	EGFR	19	TGTCGATGGACTTC CAGAAC	ATTGGGACAGCTT GGATCA	CACCTGGGCAGCTG CCAA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
EGLN3.1	NM_022073	NM_022073.2	EGLN3	2970	GCTGGTCCTCTACT GCGG	CCACCATTCGCTT AGACCTC	CCGGCTGGGCAAAT ACTACGTCAA
EGR1.1	NM_001964	NM_001964.2	EGR1	2615	GTCCCCGCTGCAG ATCTCT	CTCCAGCTTAGG GTAGTTGTCCAT	CGGATCCTTTCCTCA CTCGCCCA
EIF2C1.1	NM_012199	NM_012199.2	EIF2C1	6454	CCCTCACGGACTCT CAGC	TGGGTGACTTCC ACCTTCA	CGTTCGCTTCACCAA GGAGATCAA
EIF4EBP1.1	NM_004095	NM_004095.2	EIF4EBP1	4275	GGCGGTGAAGAGT CACAGT	TTGGTAGTGCTCC ACACGAT	TGAGATGGACATTTA AAGCACCAGCC
ELTD1.1	NM_022159	NM_022159.3	ELTD1	6154	AGGTCTTGTGCAAG AGGAGC	AACCCCAAAGATC CAGGTG	CTCGCTCTTCTGTTC CTTCTCGGC
EMCN.1	NM_016242	NM_016242.2	EMCN	3875	AGGCACTGAGGGT GGAAA	CACCGGCAAAATA ATACTGGA	AATGCAAGCACTTCA GCAACCAGC
EMP1.1	NM_001423	NM_001423.1	EMP1	986	GCTAGTACTTTGAT GCTCCCTTGAT	GAACAGCTGGAG GCCAAGTC	CCAGAGAGCCTCCC TGCAGCCA
ENO2.1	NM_001975	NM_001975.2	ENO2	6155	TCCTTGGCTTACCT GACCTC	AACCCCAATGAGT AGGGCA	CTGTCTCTGCTCGCC CTCCTTTCT
ENPEP.1	NM_001977	NM_001977.3	ENPEP	6156	CACCTACACGGAGA ACGGAC	CCTGGCATCTGTT GGTTCA	TCAAGAGCATAGTG GCCACCGATC
ENPP2.1	NM_006209	NM_006209.3	ENPP2	6174	CTCCTGCGCACTAA TACCTTC	TCCCTGGATAATT GGGTCTG	TAAGTTCCTCTGGCA TGGTTGGCC
EPAS1.1	NM_001430	NM_001430.3	EPAS1	2754	AAGCCTTGGAGGG TTTCATTG	TGCTGATGTTTTT TGACAGAAAGAT	TGTCGCCATCTTGG GTCACCACG
EPB41L3.1	NM_012307	NM_012307.2	EPB41L3	4554	TCAGTGCCATACGC TCTCAC	CTTGGGCTCCAG GTAGCA	CTCTCCTTCCCTCTG GCTCTGTGC
EPHA2.1	NM_004431	NM_004431.2	EPHA2	2297	CGCCTGTTCACCAA GATTGAC	GTGGCGTGCCTC GAAGTC	TGCGCCCGATGAGA TCACCG
EPHB1.3	NM_004441	NM_004441.3	EPHB1	6508	CCTTGGGAGGGAA GATCC	GAAGTGAAGTTGC GGTAGGC	ATGGCCTCTGGAGC TGTCCATCTC
EPHB2.1	NM_004442	NM_004442.4	EPHB2	2967	CAACCAGGCAGCT CCATC	GTAATGCTGTCCA CGGTGC	CACCTGATGCATGAT GGACACTGC
EPHB4.1	NM_004444	NM_004444.3	EPHB4	2620	TGAACGGGGTATC CTCCTTA	AGGTACCTCTCG GTCAGTGG	CGTCCCATTGAGCC TGTCATGT
EPO.1	NM_000799	NM_000799.2	EPO	5992	CAGTGCCAGCAAT GACATCT	CAAGTTGGCCCT GTGACAT	CTCTCTGGACAGTTC CTCTGGCCC
ErbB3.1	NM_001982	NM_001982.1	ERBB3	93	CGGTTATGTCATGC CAGATACAC	GAAGTGAAGACC ACTGAAGAAAGG	CCTCAAAGGTACTCC CTCCTCCCGG

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
ERBB4.3	NM_005235	NM_005235.1	ERBB4	407	TGGCTCTTAATCAG TTTCGTTACCT	CAAGGCATATCGA TCCTCATAAAGT	TGTCCCACGAATAAT GCGTAAATTCTCCAG
ERCC1.2	NM_001983	NM_001983.1	ERCC1	869	GTCCAGGTGGATG TGAAAGA	CGGCCAGGATAC ACATCTTA	CAGCAGGCCCTCAA GGAGCTG
ERCC4.1	NM_005236	NM_005236.1	ERCC4	5238	CTGCTGGAGTACG AGCGAC	GGGCGCACACTA CTAGCC	CTGGTGCTGGAAC GCTCGACACT
EREG.1	NM_001432	NM_001432.1	EREG	309	ATAACAAAGTGTAG CTCTGACATGAATG	CACACCTGCAGTA GTTTTGACTCA	TTGTTTGCATGGACA GTGCATCTATCTGGT
ERG.1	NM_004449	NM_004449.3	ERG	3884	CCAACACTAGGCTC CCCA	CCTCCGCCAGGT CTTTAGT	AGCCATATGCCTTCT CATCTGGGC
ERK1.3	NM_002746	Z11696.1	MAPK3	548	ACGGATCACAGTG GAGGAAG	CTCATCCGTCGG GTCATAGT	CGCTGGCTCACCCC TACCTG
ERK2.3	NM_002745	NM_002745.1	MAPK1	557	AGTTCTTGACCCCT GGTCCT	AAACGGCTCAAAG GAGTCAA	TCTCCAGCCCGTCTT GGCTT
ESPL1.3	NM_012291	NM_012291.1	ESPL1	2053	ACCCCCAGACCGG ATCAG	TGTAGGGCAGAC TTCCTCAAACA	CTGGCCCTCATGTC CCCTTCACG
ESRRG.3	NM_001438	NM_001438.1	ESRRG	2225	CCAGCACCATTGTT GAAGAT	AGTCTCTTGGGC ATCGAGTT	CCCCAGACCAAGTG TGAATACATGCT
F2.1	NM_000506	NM_000506.2	F2	2877	GCTGCATGTCTGGA AGGTAAGT	CCTGACCGGGTG ATGTTAC	CCTCGGTAGTTCGTA CCCAGACCCTCAG
F3.1	NM_001993	NM_001993.2	F3	2871	GTGAAGGATGTGAA GCAGACGTA	AACCGGTGCTCT CCACATTC	TGGCACGGGTCTTC TCCTACC
FABP1.1	NM_001443	NM_001443.1	FABP1	6175	GGGTCCAAAGTGAT CCAAAA	CCCTGTCATTGTC TCCAGC	ACATTCCTCCCCAC CGTGAATTC
FABP7.1	NM_001446	NM_001446.3	FABP7	4048	GGAGACAAAGTGG TCATCAGG	CTCTTCTCCCAGC TGGAAGT	TCTCAGCACATTCAA GAACACGGAGA
FAP.1	NM_004460	NM_004460.2	FAP	3403	CTGACCAGAACCAC GGCT	GGAAGTGGGTCA TGTGGG	CGGCCTGTCCACGA ACCACTTATA
fas.1	NM_000043	NM_000043.1	FAS	42	GGATTGCTCAACAA CCATGCT	GGCATTAAACTT TTGGACGATAA	TCTGGACCCTCCTAC CTCTGGTTCTTACGT
fasl.2	NM_000639	NM_000639.1	FASLG	94	GCACTTTGGGATTC TTTCCATTAT	GCATGTAAGAAGA CCCTCACTGAA	ACAACATTCTCGGTG CCTGTAACAAAGAA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
FBXW7.1	NM_033632	NM_033632.1	FBXW7	2644	CCCCAGTTTCAACG AGACTT	GTTCCAGGAATGA AAGCACA	TCATTGCTCCCTAAA GAGTTGGCACTC
FCER1G.2	NM_004106	NM_004106.1	FCER1G	4073	TGCCATCCTGTTTC TGTATGGA	TGCCTTTCGCACT TGGATCT	TTGTCCTCACCCTCC TCTACTGTCGACTG
FCGR3A.1	NM_000569	NM_000569.4	FCGR3A	3080	GTCTCCAGTGGAA GGGAAAA	AGGAATGCAGCT ACTCACTGG	CCCATGATCTTCAAG CAGGGAAGC
FDPS.1	NM_002004	NM_002004.1	FDPS	516	GGATGATTACCTTG ACCTCTTTGG	TGCATTTGTTGTC CTGGATGTC	CAGTGTGACCGGCA AAATTGGCAC
FEN1.1	NM_004111	NM_004111.4	FEN1	3938	GTGGAGAAGGGTA CGCCAG	CTCATGGCAACCA GTCCC	CGCTGAGAGACTCT GTTCTCCCTGG
FGF1.1	NM_000800	NM_000800.2	FGF1	4561	GACACCGACGGGC TTTTA	CAGCCTTTCAG GAACAAAC	ACGGCTCACAGACA CCAAATGAGG
FGF2.2	NM_002006	NM_002006.2	FGF2	681	AGATGCAGGAGAG AGGAAGC	GTTTTGCAGCCTT ACCCAAT	CCTGCAGACTGCTTT TTGCCCAAT
FGF9.1	NM_002010	NM_002010.1	FGF9	6177	CACAGCTGCCATAC TTCGAC	AAGTAAGACTGCA CCCTCGC	AGGCCACCAGCCAG AATCCTGATA
FGFR1.3	NM_023109	NM_023109.1		353	CACGGGACATTAC CACATC	GGGTGCCATCCA CTTCACA	ATAAAAAGACAACCA ACGGCCGACTGC
FGFR2 isoform 1.1	NM_000141	NM_000141.2	FGFR2	2632	GAGGGACTGTTGG CATGCA	GAGTGAGAATTC GATCCAAGTCTTC	TCCAGAGACCAAC GTTCAAGCAGTTG
FH.1	NM_000143	NM_000143.2	FH	4938	ATGGTTGCAGCCCA AGTC	CAAAATGTCCATT GCTGCC	ACAGTGACAGCAACA TGGTTCCCC
FHIT.1	NM_002012	NM_002012.1	FHIT	871	CCAGTGGAGCGCT TCCAT	CTCTCTGGGTCTG TCTGAAACAA	TCGGCCACTTCATCA GGACGCAG
FHL1.1	NM_001449	NM_001449.3	FHL1	4005	ATCCAGCCTTTGCC GAATA	CCTTGTAAGCTGG AGGGACC	TCCTATCTGCCACAC ATCCAGCGT
FIGF.1	NM_004469	NM_004469.2	FIGF	3160	GGTTCAGCTTTCT GTAGCTGT	GCCGCAGGTTCT AGTTGCT	ATTGGTGGCCACAC CACCTCCTTA
FILIP1.1	NM_015687	NM_015687.2	FILIP1	4510	ACACCGGTCACAAC GTCAT	CTGGGATGACCC GTCTTG	CCTGACACTGACTG GGTTCCTCGA
FKBP1A.1	NM_000801	NM_000801.2	FKBP1A	6330	CTGCCCTGACTGAA TGTGTT	TACGAGGAGAAA GGGGAAGA	TCACTCAGCTTTGCT TCCGACACC
FLJ22655.1	NM_024730	NM_024730.2	RERGL	3870	CTCCTTCACACAGA ACCTTTCA	AGGCAAAGTGGG ATCGCT	CACACTCACCCTAAC CTACTGGCGG
FLT1.1	NM_002019	NM_002019.3	FLT1	6062	GGCTCCTGAATCTA TCTTTG	TCCCACAGCAATA CTCCGTA	CTACAGCACCAAGA GCGACGTGTG

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
FLT3LG.1	NM_001459	NM_001459.2	FLT3LG	6178	TGGGTCCAAGATG CAAGG	GAAAGGCACATTT GGTGACA	AGTGTATCTCCGTGT TCACGCGCT
FLT4.1	NM_002020	NM_002020.1	FLT4	2782	ACCAAGAAGCTGAG GACCTG	CCTGGAAGCTGT AGCAGACA	AGCCCGCTGACCAT GGAAGATCT
FN1.1	NM_002026	NM_002026.2	FN1	4528	GGAAGTGACAGAC GTGAAGGT	ACACGGTAGCCG GTCACT	ACTCTCAGGCGGTG TCCACATGAT
FOLR1.1	NM_016730	NM_016730.1	FOLR1	859	GAACGCCAAGCAC CACAAG	CCAGGGTCGACA CTGCTCAT	AAGCCAGGCCCCGA GGACAAGTT
FOS.1	NM_005252	NM_005252.2	FOS	2418	CGAGCCCTTTGATG ACTTCCT	GGAGCGGGCTGT CTCAGA	TCCCAGCATCATCCA GGCCAG
FRAP1.1	NM_004958	NM_004958.2	MTOR	3095	AGCGCTAGAGACT GTGGACC	ATGATCCGGGAG GCATAGT	CCTGACGGAGTCCC TGGATTTTAC
FRP1.3	NM_003012	NM_003012.2	SFRP1	648	TTGGTACCTGTGG GTTAGCA	CACATCCAAATGC AAACTGG	TCCCCAGGGTAGAA TTCAATCAGAGC
FST.1	NM_006350	NM_006350.2	FST	2306	GTAAGTCGGATGA GCCTGTCTGT	CAGCTTCCTTCAT GGCACACT	CCAGTGACAATGCCA CTTATGCCAGC
FZD2.2	NM_001466	NM_001466.2	FZD2	3760	TGGATCCTCACCTG GTCCG	GCGCTGCATGTC TACCAA	TGCGCTTCACCTTC TTCAGTGTC
G-Catenin.1	NM_002230	NM_002230.1	JUP	770	TCAGCAGCAAGGG CATCAT	GGTGGTTTTCTTG AGCGTGTACT	CGCCCGCAGGCCTC ATCCT
GADD45B.1	NM_015675	NM_015675.1	GADD45B	2481	ACCCTCGACAAGAC CACACT	TGGGAGTTCATG GGTACAGA	AACTTCAGCCCCAGC TCCCAAGTC
GAS2.1	NM_005256	NM_005256.2	GAS2	6451	AACATGTCATGGTC CGTGTG	GGGGTCGTGTTT CAACAAAT	CCTGCAAAAGTTTCC CAGCCTCCT
GATA3.3	NM_002051	NM_002051.1	GATA3	1	CAAAGGAGCTCACT GTGGTGTCT	GAGTCAGAATGG CTTATTCACAGAT G	TGTTCCAACCACTGA ATCTGGACC
GATM.1	NM_001482	NM_001482.2	GATM	6296	GATCTCGGCTTGGA CGAAC	GTAGCTGCCTGG GTGCTCT	AAAGTTCGCTGCACC CATCCTGTC
GBL.1	NM_022372	NM_022372.3	MLST8	6302	GCTGTCAATAGCAC CGGAA	GGTCACCTCGTC ACCAATG	CCCCCGTCAGATTC CAGACATAGC
GBP2.2	NM_004120	NM_004120.2	GBP2	2060	GCATGGGAACCAT CAACCA	TGAGGAGTTTGC CTTGATTCTG	CCATGGACCAACTTC ACTATGTGACAGAGC
GCLC.3	NM_001498	NM_001498.1	GCLC	330	CTGTTGCAGGAAG GCATTGA	GTCAGTGGGTCT CTAATAAAGAGAT GAG	CATCTCCTGGCCCA GCATGTT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
GCLM.2	NM_002061	NM_002061.1	GCLM	704	TGTAGAATCAAAC CTTCATCATCAACT AG	CACAGAATCCAGC TGTGCAACT	TGCAGTTGACATGG CCTGTTCAAGTCC
GFRA1.1	NM_005264	NM_005264.3	GFRA1	6179	TCCGGGTAAAGAAC AAGCC	GTGGCAAAACAT GAGTGGG	TTTCATTCTCAGACC CTGCTGGCC
GJA1.1	NM_000165	NM_000165.2	GJA1	2600	GTTCACTGGGGGT GTATGG	AAATACCAACATG CACCTCTCTT	ATCCCCTCCCTCTCC ACCCATCTA
GLYAT.1	NM_201648	NM_201648.2	GLYAT	6180	TACCATTGCAAGGT GCC	GGATGCTGGGAG GCTCTT	AGGATTTCTCCAGCA TCTGCAGCA
GMNN.1	NM_015895	NM_015895.3	GMNN	3880	GTTGCTACGAGG ATTGAGC	TGCGTACCCACTT CCTGC	CCTCTTGCCCACTTA CTGGGTGGA
GNAS.1	NM_000516	NM_000516.3	GNAS	2665	GAACGTGCCTGACT TTGACTT	ACTCCTTCATCCT CCCACAG	CCTCCCGAATTCTAT GAGCATGCC
GPC3.1	NM_004484	NM_004484.2	GPC3	659	TGATGCGCCTGGA AACAGT	CGAGGTTGTGAA AGGTGCTTATC	AGCAGGCAACTCCG AAGGACAACG
GPX1.2	NM_000581	NM_000581.2	GPX1	2955	GCTTATGACCGACC CCAA	AAAGTTCCAGGCA ACATCGT	CTCATCACCTGGTCT CCGGTGTGT
GPX2.2	NM_002083	NM_002083.1	GPX2	890	CACACAGATCTCCT ACTCCATCCA	GGTCCAGCAGTG TCTCCTGAA	CATGCTGCATCCTAA GGCTCCTCAGG
GPX3.1	NM_002084	NM_002084.3	GPX3	6271	GCTCTAGGTCCAAT TGTTCTGC	TGGAGGCAGTGG GAGATG	ACTGATACCTCAACC TTGGGGCCA
GRB14.1	NM_004490	NM_004490.1	GRB14	2784	TCCCACTGAAGCCC TTTCAG	AGTGCCAGGCG TAAACATC	CCTCCAAGCGAGTC CTTCTTCAACCG
GRB7.2	NM_005310	NM_005310.1	GRB7	20	CCATCTGCATCCAT CTTGTT	GGCCACCAGGGT ATTATCTG	CTCCCCACCCTTGA GAAGTGCCT
GRO1.2	NM_001511	NM_001511.1	CXCL1	86	CGAAAAGATGCTGA ACAGTGACA	TCAGGAACAGCC ACCAGTGA	CTTCCTCCTCCCTTC TGGTCAGTTGGAT
GSTM1.1	NM_000561	NM_000561.1	GSTM1	727	AAGCTATGAGGAAA AGAAGTACACGAT	GGCCCAGCTTGA ATTTTCA	TCAGCCACTGGCTTC TGTCATAATCAGGAG
GSTM3.2	NM_000849	NM_000849.3	GSTM3	731	CAATGCCATCTTGC GCTACAT	GTCCACTCGAATC TTTTCTTCTTCA	CTCGCAAGCACAACA TGTGTGGTGAGA
GSTp.3	NM_000852	NM_000852.2	GSTP1	66	GAGACCCTGCTGT CCCAGAA	GGTTGTAGTCAG CGAAGGAGATC	TCCCACAATGAAGGT CTTGCTCCCT
GSTT1.3	NM_000853	NM_000853.1	GSTT1	813	CACCATCCCCACCC TGTCT	GGCCTCAGTGTG CATCATTCT	CACAGCCGCCTGAA AGCCACAAT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
GZMA.1	NM_006144	NM_006144.2	GZMA	4111	GAAAGAGTTTCCCT ATCCATGC	TGCTTTTTCGTC AGCTGTAA	AGCCACACGCGAAG GTGACCTTAA
HADH.1	NM_005327	NM_005327.2	HADH	6181	CCACCAGACAAGAC CGATTC	CCACAAGTTTCAT GACAGGC	CTGGCCTCCATTCT TCAACCCAG
HAVCR1.1	NM_012206	NM_012206.2	HAVCR1	6284	CCACCCAAGGTCAC GACTAC	GAACAGTGGTGC TCGTTTCG	TCACAACCTGTTCCAA CCGTCAACGA
HDAC1.1	NM_004964	NM_004964.2	HDAC1	2602	CAAGTACCACAGCG ATGACTACATTAA	GCTTGCTGTACTC CGACATGTT	TTCTTGCGCTCCATC CGTCCAGA
Hepsin.1	NM_002151	NM_002151.1	HPN	814	AGGCTGCTGGAGG TCATCTC	CTTCCTGCGGCC ACAGTCT	CCAGAGGCCGTTTC TTGGCCG
HER2.3	NM_004448	NM_004448.1	ERBB2	13	CGGTGTGAGAAGT GCAGCAA	CCTCTCGCAAGT GCTCCAT	CCAGACCATAGCACA CTCGGGCAC
HGD.1	NM_000187	NM_000187.2	HGD	6303	CTCAGGTCTGCCC CTACAAT	TTATTGGTGCTCC GTGGAC	CTGAGCAGCTCTCA GGATCGGCTT
HGF.4	M29145	M29145.1		457	CCGAAATCCAGATG ATGATG	CCCAAGGAATGA GTGGATTT	CTCATGGACCCTGG TGCTACACG
HGFAC.1	NM_001528	NM_001528.2	HGFAC	2704	CAGGACACAAGTG CCAGATT	GCAGGGAGCTGG AGTAGC	CGCTCACGTTCTCAT CCAAGTGG
HIF1A.3	NM_001530	NM_001530.1	HIF1A	399	TGAACATAAAGTCT GCAACATGGA	TGAGGTTGGTTAC TGTTGGTATCATA TA	TTGCACTGCACAGG CCACATTAC
HIF1AN.1	NM_017902	NM_017902.2	HIF1AN	7211	TGTTGGCCAGGTCT CACTG	GCATCATAGGGC CTGGAG	CTCTAGCCAGTTAGC CTCGGGCAG
HIST1H1D.1	NM_005320	NM_005320.2	HIST1H1D	4013	AAAAAGGCGAAGAA GGCAG	GCTCAGATACTG GGGGTCC	AACTGCTGGGAAAC GCAAAGCATC
HLA-B.1	NM_005514	NM_005514.6	HLA-B	6334	CTTGTGAGGGACT GAGATGC	TGCAGAAAGAGAT GCCAGAG	TCTTCACGCCTCCCC TTTGTGA
HLA-DPA1.1	NM_033554	NM_033554.2	HLA-DPA1	6314	CGCCCTGAAGACA GAATGT	TCGGAGACTCAG CAGGAAA	TGATCTTGAGAGCCC TCTCCTTGGC
HLA-DPB1.1	NM_002121	NM_002121.4	HLA-DPB1	1740	TCCATGATGGTTCT GCAGGTT	TGAGCAGCACCA TCAGTAACG	CCCCGGACAGTGGC TCTGACG
HLA-DQB1.1	NM_002123	NM_002123.3	HLA-DQB1	6304	GGTCTGCTCGGTG ACAGATT	CTCCTGATCATTG CGAAACC	TATCCAGGCCAGATC AAAGTCCGG
HLADQA1.2	NM_002122	NM_002122.3	HLA-DQA1	4071	CATCTTTCCTCCTG TGGTCA	GCTGGTCTCAGA AACACCTTC	TGTGACTGACTGCC CATTGCTCAG
HMGB1.1	NM_002128	NM_002128.3	HMGB1	2162	TGGCCTGTCCATTG GTGAT	GCTTGTCATCTGC AGCAGTGTT	TTCCACATCTCTCCC AGTTTCTTCGCAA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
HNRNPAB.1	NM_004499	NM_004499.3	HNRNPAB	6051	AGCAGGAGCGACC AACTGA	GTTTGCCAAGTTA AATTTGGTACATA AT	CTCCATATCCAAACA AAGCATGTGTGCG
HPCAL1.1	NM_002149	NM_002149.2	HPCAL1	6182	CAGGCAGATGGAC ACCAA	GTCGCTCTTGGC ACCTCT	TCTTCCAAGGACAGT TTGCCGTCA
HPD.1	NM_002150	NM_002150.2	HPD	6183	AGCTGAAGACGGC CAAGAT	CGTCGTAGTCCA CCAGGATT	AGCTCCTCCAGGGC ATCAATGTTC
HSD11B2.1	NM_000196	NM_000196.3	HSD11B2	6185	CCAACCTGCCTCAA GAGC	GGAAGTGGCCAT GCAAGT	CTGCAGGCCTACGG CAAGGACTAC
HSP90AB1.1	NM_007355	NM_007355.2	HSP90AB1	5456	GCATTGTGACCAGC ACCTAC	GAAGTGCCTGGG CTTTCAT	ATCCGCTCCATATTG GCTGTCCAG
HSPA1A.1	NM_005345	NM_005345.4	HSPA1A	2412	CTGCTGCGACAGT CCACTA	CAGGTTCGCTCT GGGAAG	AGAGTGAAGTCCCGTT GTCCCAAGG
HSPA8.1	NM_006597	NM_006597.3	HSPA8	2563	CCTCCCTCTGGTG GTGCTT	GCTACATCTACAC TTGGTTGGCTTAA	CTCAGGGCCCCACCA TTGAAGAGGTTG
HSPB1.1	NM_001540	NM_001540.2	HSPB1	2416	CCGACTGGAGGAG CATAAA	ATGCTGGCTGAC TCTGCTC	CGCACTTTTCTGAGC AGACGTCCA
HSPG2.1	NM_005529	NM_005529.2	HSPG2	1783	GAGTACGTGTGCC GAGTGTT	CTCAATGGTGACC AGGACA	CAGCTCCGTGCCTC TAGAGGCCT
HTATIP.1	NM_006388	NM_006388.2	KAT5	3893	TCGAATTGTTTGGG CACTG	GCGTGGTGCTGA CGGTAT	TGAGGACTCCCAGG ACAGCTCTGA
HYAL1.1	NM_153281	NM_153281.1	HYAL1	4524	TGGCTGTGGAGTT CAAATGT	CCAATCACCACAT GCTCTTC	CGATGCTACCCTGG CTGGCAG
HYAL2.1	NM_033158	NM_033158.2	HYAL2	5192	CAACCATGCACTCC CAGTC	ACTAAGCCCCGT GAGCCT	TCTTCACACGACCCA CCTACAGCC
HYAL3.1	NM_003549	NM_003549.2	HYAL3	6298	TATGTCCGCCTCAC ACACC	CAATGGAAGTCA CAAGGTCA	TGGGACAGGAACCT CCCAGATCTC
ICAM1.1	NM_000201	NM_000201.1	ICAM1	1761	GCAGACAGTGACC ATCTACAGCTT	CTTCTGAGACCTC TGGCTTCGT	CCGGCGCCCAACGT GATTCT
ICAM2.1	NM_000873	NM_000873.2	ICAM2	2472	GGTCATCCTGACAC TGCAAC	TGCACTCAATGGT GAAGGAC	TTGCCCACAGCCAC CAAAGTG
ICAM3.1	NM_002162	NM_002162.3	ICAM3	7219	GCCTTCAATCTCAG CAACG	GAGAGCCATTGC AGTACAC	CAGAGGATCCGACT GTTGCCAGTC
ID1.1	NM_002165	NM_002165.1	ID1	354	AGAACCGCAAGGT GAGCAA	TCCAAGTGAAGGT CCCTGATG	TGGAGATTCTCCAGC ACGTCATCGAC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
ID2.4	NM_002166	NM_002166.1	ID2	37	AACGACTGCTACTC CAAGCTCAA	GGATTTCCATCTT GCTCACCTT	TGCCCAGCATCCCC CAGAACAA
ID3.1	NM_002167	NM_002167.3	ID3	6052	CTTCACCAAATCCC TTCCTG	CTCTGGCTCTTCA GGCCACA	TCACAGTCCTTCGCT CCTGAGCAC
IFI27.1	NM_005532	NM_005532.2	IFI27	2770	CTCTCCGGATTGAC CAAGTT	TAGAACCTCGCAA TGACAGC	CAGACCCAATGGAG CCCAGGAT
IGF1.2	NM_000618	NM_000618.1	IGF1	60	TCCGGAGCTGTGAT CTAAGGA	CGGACAGAGCGA GCTGACTT	TGTATTGCGCACCCC TCAAGCCTG
IGF1R.3	NM_000875	NM_000875.2	IGF1R	413	GCATGGTAGCCGA AGATTTCA	TTTCCGGTAATAG TCTGTCTCATAGA TATC	CGCGTCATACCAAAA TCTCCGATTTTGA
IGF2.2	NM_000612	NM_000612.2	IGF2	166	CCGTGCTTCCGGA CAACTT	TGGACTGCTTCCA GGTGTC	TACCCCGTGGGCAA GTTCTTCCAA
IGFBP2.1	NM_000597	NM_000597.1	IGFBP2	373	GTGGACAGCACCA TGAACA	CCTTCATACCCGA CTTGAGG	CTTCCGGCCAGCAC TGCCTC
IGFBP3.1	NM_000598	NM_000598.4	IGFBP3	6657	ACATCCCAACGCAT GCTC	CCACGCCCTTGTT TCAGA	ACACCACAGAAGGC TGTGAGCTCC
IGFBP5.1	NM_000599	NM_000599.1	IGFBP5	594	TGGACAAGTACGG GATGAAGCT	CGAAGGTGTGGC ACTGAAAGT	CCCGTCAACGTACTC CATGCCTGG
IGFBP6.1	NM_002178	NM_002178.1	IGFBP6	836	TGAACCGCAGAGA CCAACAG	GTCTTGGACACC CGCAGAAT	ATCCAGGCACCTCTA CCACGCCCTC
IL-7.1	NM_000880	NM_000880.2	IL7	2084	GCGGTGATTTCGGA AATTCTG	CTCTCCTGGGCA CCTGCTT	CTCTGGTCCTCATCC AGGTGCGC
IL-8.1	NM_000584	NM_000584.2	IL8	2087	AAGGAACCATCTCA CTGTGTGTAAAC	ATCAGGAAGGCT GCCAAGAG	TGACTTCCAAGCTGG CCGTGGC
IL10.3	NM_000572	NM_000572.1	IL10	909	GGCGCTGTCATCG ATTTCTT	TGGAGCTTATTAA AGGCATTCTTCA	CTGCTCCACGGCCT TGCTCTTG
IL11.2	NM_000641	NM_000641.2	IL11	2166	TGGAAGGTTCCACA AGTCAC	TCTTGACCTTGCA GCTTTGT	CCTGTGATCAACAGT ACCCGTATGGG
IL15.1	NM_000585	NM_000585.2	IL15	6187	GGCTGGGTACCAA TGCTG	TGAGAGCCAGTA GTCAGTGGTT	CAGCTATGCTGGTA GGCTCCTGCC
IL1B.1	NM_000576	NM_000576.2	IL1B	2755	AGCTGAGGAAGAT GCTGGTT	GGAAAGAAGGTG CTCAGGTC	TGCCACAGACCTTC CAGGAGAAT
IL6.3	NM_000600	NM_000600.1	IL6	324	CCTGAACCTTCCAA AGATGG	ACCAGGCAAGTC TCCTCATT	CCAGATTGGAAGCAT CCATCTTTTCA
IL6ST.3	NM_002184	NM_002184.2	IL6ST	2317	GGCCTAATGTTCCA GATCCT	AAAATTGTGCCTT GGAGGAG	CATATTGCCAGTGG TCACCTCACA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
ILT-2.2	NM_006669	NM_006669.1	LILRB1	583	AGCCATCACTCTCA GTGCAG	ACTGCAGAGTCA GGGTCTCC	CAGGTCCTATCGTG GCCCCTGA
IMP3.1	NM_018285	NM_018285.2	IMP3	4751	GTGGACTCGTCCAA GATCAA	GGCTTCCAGATC GAAGTCAT	CTCATTGTACTCTAG CACGTGCCGC
INDO.1	NM_002164	NM_002164.3	IDO1	5124	CGCCTTGACAGTCT AGTTC	ATCTCCATGACCT TTGCC	ACATATGCCATGGTG ATGCATCCC
INHBA.1	NM_002192	NM_002192.1	INHBA	2635	GTGCCCGAGCCAT ATAGCA	CGGTAGTGGTTG ATGACTGTTGA	ACGTCCGGGTCTCTC ACTGTCCTTCC
INHBB.1	NM_002193	NM_002193.1	INHBB	2636	AGCCTCCAGGATAC CAGCAA	TCTCCGACTGACA GGCATTG	AGCTAAGCTGCCATT TGTCACCG
INSR.1	NM_001079817	NM_001079817.1	INSR	6455	CAGTCTCCGAGAG CGGATT	GTGATGGCAGGT GAAGCC	AGTTCCTCAATGAGG CCTCGGTCA
IQGAP2.1	NM_006633	NM_006633.2	IQGAP2	6453	AGAGACACCAGCAA CTGCG	ATCATTGCACGGC TCACC	CCGTGGCATGGTCT ACCTCCTGTT
ISG20.1	NM_002201	NM_002201.4	ISG20	6189	GTGTCAGACTGAAG CCCAT	GTTGCTGTCCCAA AAAGCC	AAAGCCTCTAGTCCC TGCGGAACG
ITGA3.2	NM_002204	NM_002204.1	ITGA3	840	CCATGATCCTCACT CTGCTG	GAAGCTTTGTAGC CGGTGAT	CACTCCAGACCTCG CTTAGCATGG
ITGA4.2	NM_000885	NM_000885.2	ITGA4	2867	CAACGCTTCAGTGA TCAATCC	GTCTGGCCGGGA TTCTTT	CGATCCTGCATCTGT AAATCGCCC
ITGA5.1	NM_002205	NM_002205.1	ITGA5	2668	AGGCCAGCCCTAC ATTATCA	GTCTTCTCCACAG TCCAGCA	TCTGAGCCTTGTCTT CTATCCGGC
ITGA6.2	NM_000210	NM_000210.1	ITGA6	2791	CAGTGACAAACAGC CCTTCC	GTTTAGCCTCATG GGCGTC	TCGCCATCTTTTGTG GGATTCTT
ITGA7.1	NM_002206	NM_002206.1	ITGA7	259	GATATGATTGGTCG CTGCTTTG	AGAACTTCCATT CCCACCAT	CAGCCAGGACCTGG CCATCCG
ITGAV.1	NM_002210	NM_002210.2	ITGAV	2671	ACTCGGACTGCACA AGCTATT	TGCCATCACCATT GAAATCT	CCGACAGCCACAGA ATAACCCAAA
ITGB1.1	NM_002211	NM_002211.2	ITGB1	2669	TCAGAATTGGATTT GGCTCA	CCTGAGCTTAGCT GGTGTTG	TGCTAATGTAAGGCA TCACAGTCTTTTCCA
ITGB3.1	NM_000212	NM_000212.2	ITGB3	6056	ACCGGGGAGCCCT ACATGA	CCTTAAGCTCTTT CACTGACTCAATC T	AAATACCTGCAACCG TTACTGCOGTGAC
ITGB4.2	NM_000213	NM_000213.2	ITGB4	2793	CAAGGTGCCCTCA GTGGA	GCGCACACCTTC ATCTCAT	CACCAACCTGTACCC GTATTGCGA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
ITGB5.1	NM_002213	NM_002213.3	ITGB5	2670	TCGTGAAAGATGAC CAGGAG	GGTGAACATCATG ACGCAGT	TGCTATGTTTCTACA AAACCGCCAAGG
JAG1.1	NM_000214	NM_000214.1	JAG1	4190	TGGCTTACACTGGC AATGG	GCATAGCTGTGA GATGCGG	ACTCGATTTCCAGC CAACCACAG
K-ras.10	NM_033360	NM_033360.2	KRAS	3090	GTCAAAATGGGGA GGGACTA	CAGGACCACCAC AGAGTGAG	TGTATCTTGTTGAGC TATCCAAACTGCCC
KCNJ15.1	NM_002243	NM_002243.3	KCNJ15	6299	GGACGTTCTACCTG CCTTGA	AGGCTCTGGAAA CACTGGTC	TCACTCCGCAGGTC AGGTGTCTTC
KDR.6	NM_002253	NM_002253.1	KDR	463	GAGGACGAAGGCC TCTACAC	AAAAATGCCTCCA CTTTTGC	CAGGCATGCAGTGT TCTTGGCTGT
Ki-67.2	NM_002417	NM_002417.1	MKI67	145	CGGACTTTGGGTG CGACTT	TTACAACTCTTCC ACTGGGACGAT	CCACTTGTGGAACCA CCGCTCGT
KIAA1303 raptor.1	NM_020761	NM_020761.2	RPTOR	6300	ACTACAGCGGGAG CAGGAG	GGCATCTGAGCA AGAGGGT	TGGAGGTAGCTGCA ATCAACCCAA
KIF1A.1	NM_004321	NM_004321.4	KIF1A	4015	CTCCTACTGGTCGC ACACC	TCCCGGTACACCT GCTTC	CCTGAGGACATCAAC TACGCGTCG
Kitlmg.4	NM_000899	NM_000899.1	KITLG	68	GTCCCCGGGATGG ATGTT	GATCAGTCAAGCT GTCTGACAATTG	CATCTCGCTTATCCA ACAATGACTTGGA
KL.1	NM_004795	NM_004795.2	KL	6191	GAGGTCCTGTCTAA ACCCTGTG	CTATGTGCAAGG CCCTCAA	CCTGAGGGATCTGT CTCACTGGCA
KLK3.1	NM_001648	NM_001648.2	KLK3	4172	CCAAGCTTACCACC TGCAC	AGGGTGAGGAAG ACAACCG	ACCCACATGGTGACA CAGCTCTCC
KLRK1.2	NM_007360	NM_007360.1	KLRK1	3805	TGAGAGCCAGGCT TCTTGTA	ATCCTGGTCCTCT TTGCTGT	TGTCTCAAATGCCA GCCTTCTGAA
KRT19.3	NM_002276	NM_002276.1	KRT19	521	TGAGCGGCAGAAT CAGGAGTA	TGCGGTAGGTGG CAATCTC	CTCATGGACATCAAG TCGCGGCTG
KRT5.3	NM_000424	NM_000424.2	KRT5	58	TCAGTGGAAGAAG AGTTGGA	TGCCATATCCAGA GGAAACA	CCAGTCAACATCTCT GTTGTCACAAGCA
KRT7.1	NM_005556	NM_005556.3	KRT7	4016	TTCAGAGATGAACC GGGC	ACTTGGCACGCT GGTTCT	ATGTTGTGATCTCA GCCTGCAGC
L1CAM.1	NM_000425	NM_000425.2	L1CAM	4096	CTTGCTGGCCAATG CCTA	TGATTGTCCGCA GTCAGG	ATCTACGTTGTCCAG CTGCCAGCC
LAMA3.1	NM_000227	NM_000227.2	LAMA3	2529	CAGATGAGGCACAT GGAGAC	TTGAAATGGCAGA ACGGTAG	CTGATTCCTCAGGTC CTTGGCCTG
LAMA4.1	NM_002290	NM_002290.3	LAMA4	5990	GATGCACTGCGGTT AGCAG	CAGAGGATACGC TCAGCACC	CTCTCCATCGAGGAA GGCAAATCC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
LAMB1.1	NM_002291	NM_002291.1	LAMB1	3894	CAAGGAGACTGGG AGGTGTC	CGGCAGAACTGA CAGTGTTC	CAAGTGCCTGTACCA CACGGAAGG
LAMB3.1	NM_000228	NM_000228.1	LAMB3	2530	ACTGACCAAGCCTG AGACCT	GTCACACTTGCAG CATTTCA	CCACTCGCCATACTG GGTGCAGT
LAMC2.2	NM_005562	NM_005562.1	LAMC2	997	ACTCAAGCGGAAAT TGAAGCA	ACTCCCTGAAGC CGAGACACT	AGGTCTTATCAGCAC AGTCTCCGCCTCC
LAPTM5.1	NM_006762	NM_006762.1	LAPTM5	4017	TGCTGGACTTCTGC CTGAG	TGAGATAGGTGG GCACTTCC	TCCTGACCCTCTGCA GCTCCTACA
LDB1.2	NM_003893	NM_003893.4	LDB1	6720	AACACCCAGTTTGA CGCAG	CCAGTGCAGGGG AGTTGT	AAAGCTGTCCTCGTC GTCAATGCC
LDB2.1	NM_001290	NM_001290.2	LDB2	3871	ATCACGGTGGACT GCGAC	TACCTTGGTAAAC ATGGGCTTC	AGTGTACCATGGTCA CCCAGCACG
LDHA.2	NM_005566	NM_005566.1	LDHA	3935	AGGCTACACATCCT GGGCTA	CCCGCCTAAGATT CTTCATT	TCTGCCAAATCTGCT ACAGAGAGTCCA
LGALS1.1	NM_002305	NM_002305.3	LGALS1	6305	GGGTGGAGTCTTC TGACAGC	AGACCACAAGCC ATGATTGA	CCCGGGAACATCCT CCTGGAC
LGALS3.1	NM_002306	NM_002306.1	LGALS3	2371	AGCGGAAAATGGC AGACAAT	CTTGAGGGTTTG GGTTTCCA	ACCCAGATAACGCAT CATGGAGCGA
LGALS9.1	NM_009587	NM_009587.2	LGALS9	5458	AGTACTTCCACCGC GTGC	GACAGCTGCACA GAGCCAT	CTTCCACCGTGTGG ACACCATCTC
LIMK1.1	NM_016735	NM_016735.1		3888	GCTTCAGGTGTTGT GACTGC	AAGAGCTGCCCCA TCCTTCTC	TGCCTCCCTGTCCG ACCAGTACTA
LMNB1.1	NM_005573	NM_005573.1	LMNB1	1708	TGCAAACGCTGGT GTCACA	CCCCACGAGTTCT GGTTCTTC	CAGCCCCCCTACTG ACCTCATC
LMO2.1	NM_005574	NM_005574.2	LMO2	5346	GGCTGCCAGCAGA ACATC	CTCAGGCAGTCC TCGTGC	CGCTACTTCCTGAAG GCCATCGAC
LOX.1	NM_002317	NM_002317.3	LOX	3394	CCAATGGGAGAACAA ACGG	CGCTGAGGCTGG TACTGTG	CAGGCTCAGCAAGC TGAACACCTG
LRP2.1	NM_004525	NM_004525.1	LRP2	4112	GGCTGTAGACTGG GTTTCCA	GAGACAAAGAGG CCATCCAG	CGGGCATCCAACCA GTAGAGCTTT
LRRC2.1	NM_024512	NM_024512.2	LRRC2	6315	CCAGTGTCCCAATC TGTGTC	GGTCAGGTTATTG CTGCTGA	CCACTGCAAATTCTGA CATCCGC
LTF.1	NM_002343	NM_002343.2	LTF	6269	AACGGAAGCCTGT GACTGA	AGACACCACGGC ATGATTC	CTAGAAGCTGCCATC TTGCCATGG
LYZ.1	NM_000239	NM_000239.1	LYZ	6268	TTGCTGCAAGATAA CATCGC	ACCCATGCTCTAA TGCCTTG	CACGGACAACCCTCT TTGCACAAG

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
MADH2.1	NM_005901	NM_005901.2	SMAD2	2672	GCTGCCTTTGGTAA GAACATGTC	ATCCCAGCAGTCT CTTCACAACT	TCCATCTTGCCATTG ACGCCGC
MADH4.1	NM_005359	NM_005359.3	SMAD4	2565	GGACATTACTGGCC TGTTTACA	ACCAATACTCAGG AGCAGGATGA	TGCATTCCAGCCTCC CATTTCCTCA
MAL.1	NM_002371	NM_002371.2	MAL	6194	GTTGGGAGCTTGC TGTGTC	CACAAACAGGAG GTGACCCT	ACCTCCAACTGCTGT GCTGTCTGC
MAL2.1	NM_052886	NM_052886.1	MAL2	5113	CCTTCGTCTGCCTG GAGAT	GGAACATTGGAG GAGGCAA	CAAAATCCAGACAAG ACCCCCGAA
MAP2K1.1	NM_002755	NM_002755.2	MAP2K1	2674	GCCTTTCTTACCCA GAAGCAGAA	CAGCCCCCAGCT CACTGAT	TCTCAAAGTCGTCTAT CCTTCAGTTCTCCCA
MAP2K3.1	NM_002756	NM_002756.2	MAP2K3	4372	GCCCTCCAATGTCC TTATCA	GTAGCCACTGAT GCCAAAGTC	CACATCTTCACATGG CCCTCCTTG
MAP4.1	NM_002375	NM_002375.2	MAP4	2066	GCCGGTCAGGCAC ACAAG	GCAGCATACACAC AACAAAATGG	ACCAACCAGTCCAC GCTCCAAGGG
MARCKS.1	NM_002356	NM_002356.4	MARCKS	4021	CCCCTCTTGGATCT GTTGAG	CGGTCTTGGAGA ACTGGG	CCCATGCTGGCTTCT TCAACAAAG
Maspin.2	NM_002639	NM_002639.1	SERPINB5	362	CAGATGGCCACTTT GAGAACATT	GGCAGCATTAAC CACAAGGATT	AGCTGACAACAGTGT GAACGACCAGACC
MCAM.1	NM_006500	NM_006500.2	MCAM	3972	CGAGTTCCAGTGG CTGAGA	TGCAACTGAAGCA CAGGC	CTTTCCAGCACCTGG CCTGTCTCT
MCM2.2	NM_004526	NM_004526.1	MCM2	580	GACTTTTGCCCGCT ACCTTTC	GCCACTAACTGCT TCAGTATGAAGAG	ACAGCTCATTGTTGT CACGCCGGA
MCM3.3	NM_002388	NM_002388.2	MCM3	524	GGAGAACAATCCCC TTGAGA	ATCTCCTGGATG GTGATGGT	TGGCCTTTCTGTCTA CAAGGATCACCA
MCM6.3	NM_005915	NM_005915.2	MCM6	614	TGATGGTCCTATGT GTCACATTCA	TGGGACAGGAAA CACACCAA	CAGGTTTCATACCAA CACAGGCTTCAGCA C
MCP1.1	NM_002982	NM_002982.1	CCL2	700	CGCTCAGCCAGAT GCAATC	GCACTGAGATCTT CCTATTGGTGAA	TGCCCCAGTCACCT GCTGTTA
MDH2.1	NM_005918	NM_005918.2	MDH2	2849	CCAACACCTTTGTT GCAGAG	CAATGACAGGGA CGTTGACT	CGAGCTGGATCCAA ACCCTTCAG
MDK.1	NM_002391	NM_002391.2	MDK	3231	GGAGCCGACTGCA AGTACA	GACTTTGGTGCCT GTGCC	ATCACACGCACCCCA GTTCTCAA
MDM2.1	NM_002392	NM_002392.1	MDM2	359	CTACAGGGACGCC ATCGAA	ATCCAACCAATCA CCTGAATGTT	CTTACACCAGCATCA AGATCCGG

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
MGMT.1	NM_002412	NM_002412.1	MGMT	689	GTGAAATGAAACGC ACCACA	GACCCTGCTCAC AACCAGAC	CAGCCCTTTGGGGA AGCTGG
mGST1.2	NM_020300	NM_020300.2	MGST1	806	ACGGATCTACCACA CCATTGC	TCCATATCCAACA AAAAAACTCAAAG	TTTGACACCCCTTCC CCAGCCA
MICA.1	NM_000247	NM_000247.1	MICA	5449	ATGGTGAATGTCAC CCGC	AAGCCAGAAGCC CTGCAT	CGAGGCCTCAGAGG GCAACATTAC
MIF.2	NM_002415	NM_002415.1	MIF	3907	CCGGACAGGGTCT ACATCA	GGTGGAGTTGTT CCAGCC	CTATTACGACATGAA CGCGGCCAA
MMP1.1	NM_002421	NM_002421.2	MMP1	2167	GGGAGATCATCGG GACAACTC	GGGCCTGGTTGA AAAGCAT	AGCAAGATTTCTCC AGGTCCATCAAAGG
MMP10.1	NM_002425	NM_002425.1	MMP10	4920	TGTACCCACTCTAC AACTCATTCA	TGAATGCCATTCA CATCATCTTG	AGCTCGCCCAGTTC CGCCTTTC
MMP14.1	NM_004995	NM_004995.2	MMP14	4022	GCTGTGGAGCTCT CAGGAA	AGCAAGGACAGG GACCAA	CCTGAGGAAGGCAC ACTTGCTCCT
MMP2.2	NM_004530	NM_004530.1	MMP2	672	CCATGATGGAGAG GCAGACA	GGAGTCCGTCCT TACCGTCAA	CTGGGAGCATGGCG ATGGATACCC
MMP7.1	NM_002423	NM_002423.2	MMP7	2647	GGATGGTAGCAGT CTAGGGATTA	GGAATGTCCCAT CCCAAAGAA	CCTGTATGCTGCAAC TCATGAACTTGGC
MMP9.1	NM_004994	NM_004994.1	MMP9	304	GAGAACCAATCTCA CCGACA	CACCCGAGTGTA ACCATAGC	ACAGGTATTCCTCTG CCAGCTGCC
MRP1.1	NM_004996	NM_004996.2	ABCC1	15	TCATGGTGCCCGTC AATG	CGATTGTCTTTGC TCTTCATGTG	ACCTGATACGTCTTG GTCTTCATCGCCAT
MRP2.3	NM_000392	NM_000392.1	ABCC2	55	AGGGGATGACTTG GACACAT	AAAAGTGCATGGC TTTGTC	CTGCCATTGACATG ACTGCAATTT
MRP3.1	NM_003786	NM_003786.2	ABCC3	8	TCATCCTGGCGATC TACTTCCT	CCGTTGAGTGGA ATCAGCAA	TCTGTCTGCGCTGG AGTCGCTTTCAT
MRP4.2	NM_005845	NM_005845.3	ABCC4	6057	AGCGCCTGGAATCT ACAAC	AGAGCCCCTGGA GAGAAGAT	CGGAGTCCAGTGTTT TCCCACTTA
MSH2.3	NM_000251	NM_000251.1	MSH2	2127	GATGCAGAATTGAG GCAGAC	TCTTGGAAGTC GGTTAAGA	CAAGAAGATTTACTT CGTCGATTCCCAGA
MSH3.2	NM_002439	NM_002439.1	MSH3	2132	TGATTACCATCATG GCTCAGA	CTTGTGAAAATGC CATCCAC	TCCCAATTGTGCGTT CTTCTGCAG
MSH6.3	NM_000179	NM_000179.1	MSH6	2136	TCTATTGGGGGATT GGTAGG	CAAATTGCGAGTG GTGAAAT	CCGTTACCAGCTGG AAATTCCTGAGA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
MT1B.1	NM_005947	NM_005947.1	MT1B	5355	GTGGGCTGTGCCA AGTGT	ACAGCAGCGGCA CTTCTC	ATGAGCCTTTGCAGA CACAGCCCT
MT1G.1	NM_005950	NM_005950.1	MT1G	6333	GTGCACCCACTGC CTCTT	AGCAGTTGGGGT CCATTG	CCCGAGGCGAGACT AGAGTTCCC
MT1H.1	NM_005951	NM_005951.2	MT1H	6332	CGTGTTCCTACTGCC TCTTC	AGCAGTTGGGGT CCATTG	CCGAGGTGAGACTG GAGTTCCCA
MT1X.1	NM_005952	NM_005952.2	MT1X	3897	CTCCTGCAAATGCA AAGAGTG	ACTTGGCACAGC CCACAG	CACCTCCTGCAAGAA GAGCTGCTG
MUC1.2	NM_002456	NM_002456.1	MUC1	335	GGCCAGGATCTGT GGTGGTA	CTCCACGTCGTG GACATTGA	CTCTGGCCTTCCGA GAAGGTACC
MVP.1	NM_017458	NM_017458.1	MVP	30	ACGAGAACGAGGG CATCTATGT	GCATGTAGGTGC TTCCAATCAC	CGCACCTTTCCGGT CTTGACATCCT
MX1.1	NM_002462	NM_002462.2	MX1	2706	GAAGGAATGGGAA TCAGTCATGA	GTCTATTAGAGTC AGATCCGGGACA T	TCACCCTGGAGATCA GCTCCCGA
MYBL2.1	NM_002466	NM_002466.1	MYBL2	1137	GCCGAGATCGCCA AGATG	CTTTTGATGGTAG AGTTCCAGTGATT C	CAGCATTGTCTGTCC TCCCTGGCA
MYH11.1	NM_002474	NM_002474.1	MYH11	1734	CGGTACTTCTCAGG GCTAATATATACG	CCGAGTAGATGG GCAGGTGTT	CTCTTCTGCGTGGT GGTCAACCCCTA
MYRIP.2	NM_015460	NM_015460.1	MYRIP	1704	CCTTCACCTTCCTC GTCAAC	AGCAGCTCTTGCA GACATTG	ATTTGCAATCTCCAC ACTGGCGCT
NBN.1	NM_002485	NM_002485.4	NBN	4121	GCATCTACTTGCCA GAACCAA	TCCCTTGCACT GGAGTT	CTTCCAAGTTCTGGC TGCTTGCA
NCF1.1	NM_000265	NM_000265.2	NCF1	4676	GACACCTTCATCCG TCACAT	ATAGTGCTGGCT GGGTACG	AAGCGCTTCTCAAAG CCCAGCAG
NFAT5.1	NM_006599	NM_006599.2	NFAT5	3071	CTGAACCCCTCTCC TGGTC	AGGAAACGATGG CGAGGT	CGAGAATCAGTCCC CGTGGAGTTC
NFATC2.1	NM_173091	NM_173091.2	NFATC2	5123	CAGTCAAGGTCAGA GGCTGAG	CTTTGGCTCGTG GCATTC	CGGGTTCCTACCCC ACAGTCATTC
NFKBp50.3	NM_003998	NM_003998.1	NFKB1	3439	CAGACCAAGGAGAT GGACCT	AGCTGCCAGTGC TATCCG	AAGCTGTAAACATGA GCCGCACCA
NFKBp65.3	NM_021975	NM_021975.1	RELA	39	CTGCCGGGATGGC TTCTAT	CCAGGTTCTGGA AACTGTGGAT	CTGAGCTCTGCCCG GACCGCT
NFX1.1	NM_002504	NM_002504.3	NFX1	4025	CCCTGCCATACCAG CTCA	CGTCCACATTAC ACTGTAGC	CCTGCCCTGTGACT GCTTGTAAGC

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NME2.1	NM_002512	NM_002512.2	NME2	3899	ATGCTTGGGGAGA CCAATC	CTGAATGCAGAA GTCCCCAC	AGCAGATTCAAAGCC AGGCACCAT
NNMT.1	NM_006169	NM_006169.2	NNMT	5101	CCTAGGGCAGGGA TGGAG	CTAGTCCAGCCAA ACATCCC	CCCTCTCTCATGCC CAGACTCTC
NOL3.1	NM_003946	NM_003946.3	NOL3	6307	CAGCCTTGGGAAG TGAGACT	ATGATGTGTGTG GCCTTTGT	CTCAAGGTCCCTTTC TGCTCCCT
NOS2A.3	NM_000625	NM_000625.3	NOS2	6509	GGGTCCATTATGAC TCCCAA	GCTCATCTGGAG GGGTAGG	TGTCCCTGGGTCTT CTGGTCAAAC
NOS3.1	NM_000603	NM_000603.2	NOS3	2624	ATCTCCGCCTCGCT CATG	TCGGAGCCATAC AGGATTGTC	TTCACCTCGCTTCGCC ATCACCG
NOTCH1.1	NM_017617	NM_017617.2	NOTCH1	2403	CGGGTCCACCACT TTGAATG	GTTGTATTGGTTC GGCACCAT	CCGCTCTGCAGCCG GGACA
NOTCH2.1	NM_024408	NM_024408.2	NOTCH2	2406	CACTTCCCTGCTGG GATTAT	AGTTGTCAAACAG GCACTCG	CCGTGTTGCACAGC TCATCACACT
NOTCH3.1	NM_000435	NM_000435.2	NOTCH3	6464	TGTGGACGAGTGT GCTGG	ACTCCCTGCCAG GTTGGT	ACCCTGTGGCCCTC ATGGTATCTG
NPD009 (ABAT official)	NM_020686	NM_020686.2	ABAT	1707	GGCTGTGGCTGAG GCTGTAG	GGAGCATTTCGAG GTCAAATCA	TTCCCAGAGTGTCTC ACCTCCAGCAGAG
NPM1.2	NM_002520	NM_002520.2	NPM1	2328	AATGTTGTCCAGGT TCTATTGC	CAAGCAAAGGGT GGAGTTC	AACAGGCATTTTGGGA CAACACATTCTTG
NPPB.1	NM_002521	NM_002521.2	NPPB	6196	GACACCTGCTTCTG ATTCCAC	TGAGTCACTTCAA AGGCGG	AGGGGCTTTTTCTC AACCTGTG
NPR1.1	NM_000906	NM_000906.2	NPR1	6197	ACATCTGCAGCTCC CCTG	CACACAAGCCAG CTTCCA	CCTTCAGAACCCTCA TGCTCCTGG
NPY1R.1	NM_000909	NM_000909.4	NPY1R	4513	GGATCTTCCCCACT CTGCT	TTGTCTTTTTCGC TCCTGC	CCTTCCATTCCCACC CTTCCTTCT
NRG1.3	NM_013957	NM_013957.1	NRG1	410	CGAGACTCTCCTCA TAGTGAAAGGTAT	CTTGGCGTGTGG AAATCTACAG	ATGACCACCCCGGC TCGTATGTCA
NUDT1.1	NM_002452	NM_002452.3	NUDT1	4564	ACTGGTTTCCACTC CTGCTT	GTCCAGGATGGT GTCCTGA	CCACGGGTACTTCAA GTTCCAGGG
OGG1.1	NM_002542	NM_002542.4	OGG1	6198	ACCAAGGTGGCTG ACTGC	ATATGGACATCCA CGGGC	TCTGCCTGATGGCC CTAGACAAGC
OPN, osteopontin.3	NM_000582	NM_000582.1	SPP1	764	CAACCGAAGTTTTT ACTCCAGTT	CCTCAGTCCATAA ACCACACTATCA	TCCCCACAGTAGACA CATATGATGGCCG

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
p21.3	NM_000389	NM_000389.1	CDKN1A	33	TGGAGACTCTCAG GGTCGAAA	GGCGTTTGGAGT GGTAGAAATC	CGGCGGCAGACCAG CATGAC
p27.3	NM_004064	NM_004064.1	CDKN1B	38	CGGTGGACCACGA AGAGTTAA	GGCTCGCCTCTT CCATGTC	CCGGGACTTGGAGA AGCACTGCA
P53.2	NM_000546	NM_000546.2	TP53	59	CTTTGAACCCTTGC TTGCAA	CCCGGGACAAAG CAAATG	AAGTCCTGGGTGCTT CTGACGCACA
PAH.1	NM_000277	NM_000277.1	PAH	6199	TGGCTGATTCCATT AACAGTGA	CACATTCTGTCCA TGGCTTTAC	ATCCTTTGCAGTGCC CTCCAGAAA
PAI1.3	NM_000602	NM_000602.1	SERPINE1	36	CCGCAACGTGGTTT TCTCA	TGCTGGGTTTCTC CTCCTGTT	CTCGGTGTTGGCCA TGCTCCAG
Pak1.2	NM_002576	NM_002576.3	PAK1	3421	GAGCTGTGGGTTG TTATGGA	CCATGCAAGTTTC TGTCACC	ACATCTGTCAAGGAG CCTCCAGCC
PARD6A.1	NM_016948	NM_016948.2	PARD6A	4514	GATCCTCGAGGTCA ATGGC	ACCATCATGTCCG TCACTTG	TCCAAGGTCTTCCCG GCTACTTCA
PBOV1.1	NM_021635	NM_021635.1	PBOV1	3936	GCAAAGCCTTTCCA GAAAAA	GGCTGGGCTTAA ACAGTCAT	TGGTAGCAGAATTGC CTTTTCAACCA
PCCA.1	NM_000282	NM_000282.2	PCCA	6250	GGTGAAATCTGTGC ACTGTCA	ATTCCAGCTCCAC GAGCA	TCCCCTTCTCCAAC GTGTCTCCA
PCK1.1	NM_002591	NM_002591.2	PCK1	6251	CTTAGCATGGCCCA GCAC	CTTCCGGAACCA GTTGACA	CAGCCAAACTGCC AAGATCTTCC
PCNA.2	NM_002592	NM_002592.1	PCNA	148	GAAGGTGTTGGAG GCACTCAAG	GGTTTACACCGCT GGAGCTAA	ATCCCAGCAGGCCT CGTTGATGAG
PCSK6.1	NM_002570	NM_002570.3	PCSK6	6282	ACCTTGAGTAGCAG AGGCC	GTTGCTGGAGCC ATTTCAC	CACACCTTCCTCAGA ATGGACCCC
PDCD1.1	NM_005018	NM_005018.2	PDCD1	6286	GACAACGCCACCTT CACC	GGCTCATGCGGT ACCAGT	TCTCCAACACATCGG AGAGCTTCG
PDE4DIP.1	NM_014644	NM_014644.4	PDE4DIP	6417	GCTTCGTCTTGCTG TGAGAG	AGCTTCATTGGAG GAGAGGA	TCGCGCAGTCTCTCT AAGTCATGATCTC
PDGFA.3	NM_002607	NM_002607.2	PDGFA	56	TTGTTGGTGTGCCC TGGTG	TGGGTCTGTCCA AACACTGG	TGGTGGCGGTCACT CCCTCTGC
PDGFB.3	NM_002608	NM_002608.1	PDGFB	67	ACTGAAGGAGACC CTTGGAG	TAAATAACCCTGC CCACACA	TCTCCTGCCGATGC CCCTAGG
PDGFC.3	NM_016205	NM_016205.1	PDGFC	29	AGTTACTAAAAAAT ACCACGAGGTCCTT	GTCGGTGAGTGA TTTGTGCAA	CCCTGACACCGGTC TTTGGTCTCAACT
PDGFD.2	NM_025208	NM_025208.2	PDGFD	31	TATCGAGGCAGGT CATACCA	TAACGCTTGGCAT CATCATT	TCCAGGTCAACTTTT GACTTCCGGT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
PDGFRA.2	NM_006206	NM_006206.2	PDGFRA	24	GGGAGTTTCCAAGA GATGGA	CTTCAACCACCTT CCCAAAC	CCCAAGACCCGACC AAGCACTAG
PDGFRb.3	NM_002609	NM_002609.2	PDGFRB	464	CCAGCTCTCCTTCC AGCTAC	GGGTGGCTCTCA CTTAGCTC	ATCAATGTCCCTGTC CGAGTGCTG
PDZK1.1	NM_002614	NM_002614.3	PDZK1	6319	AATGACCTCCACCT TCAACC	CGCAGGAAGAAG CCATAGTT	TGCCCTTCTTGCTTG GACAGTTTACA
PDZK3.1	NM_015022	NM_015022.2		3885	GAGCTGAGAGCCT TGAGCAT	CTCGGCCCTGCT GAGTAA	CTCGCTGCAGAGCT TGTC AAGGTC
PF4.1	NM_002619	NM_002619.1	PF4	6326	GCAGTGCCCTGTGT GTGAAG	GGCCTTGATCAC CTCCAG	TCCGTCCCAGGCAC ATCACC
PFKP.1	NM_002627	NM_002627.3	PFKP	6252	AGCTGATGCCGCAT ACATT	GGTGCTCCACGT TGGACT	CAGATCCCTGATGTC GAAGGGCTC
PFN2.1	NM_053024	NM_053024.1	PFN2	3426	TCTATACGTCGATG GTGACTGC	GCCGACAGCCAC ATTGTAT	CTCCCCACCTTGACT CTTTGTCCG
PGF.1	NM_002632	NM_002632.4	PGF	4026	GTGGTTTTCCCTCG GAGC	AGCAAGGGAACA GCCTCAT	ATCTTCTCAGACGTC CCGAGCCAG
PI3K.2	NM_002646	NM_002646.2	PIK3C2B	368	TGCTACCTGGACAG CCCG	AGGCCGTCTTC AGTAACCA	TCCTCCTGAAACGAG CTGTGTCTGACTT
PI3KC2A.1	NM_002645	NM_002645.2	PIK3C2A	6064	ATACCAATCACCGC ACAAACC	CACACTAGCATT TCTCCGCATA	TGTGCTGTGACTGG ACTTAACAAATAGCC T
PIK3CA.1	NM_006218	NM_006218.1	PIK3CA	2962	GTGATTGAAGAGCA TGCCAA	GTCCTGCGTGCG AATAGC	TCCTGCTTCTCGGGA TACAGACCA
PLA2G4C.1	NM_003706	NM_003706.1	PLA2G4C	6249	CCCTTTCCCCAAGT AGAAGAG	AGGATGTAGCAG CTGGCG	CCTTGGACCACAAAT CCAGCTCAG
PLAT.1	NM_033011	NM_033011.2	PLAT	6459	GATTTGCTGGGAAG TGCTGT	TAGCTGATGCCCT GGTCC	TAGATACCAGGGCC ACGTGCTACG
PLAUR.3	NM_002659	NM_002659.1	PLAUR	708	CCCATGGATGCTCC TCTGAA	CCGGTGGCTACC AGACATTG	CATTGACTGCCGAG GCCCCATG
PLG.1	NM_000301	NM_000301.1	PLG	6310	GGCAAAATTTCCAA GACCAT	ATGTATCCATGAG CGTGTGG	TGCCAGGCCTGGGA CTCTCA
PLN.1	NM_002667	NM_002667.2	PLN	3886	TGATGCTTCTCTGA AGTTCTGC	CCTGTCTGCATG GGATGAC	AGATCTGCAGCTTGC CACATCAGC
PLOD2.1	NM_000935	NM_000935.2	PLOD2	3820	CAGGGAGGTGGTT GCAAT	TCTCCAGGATG CATGAAG	TCCAGCCTTTTCGTG GTGACTCAA
PLP1.1	NM_000533	NM_000533.3	PLP1	4027	AGAACAGACTGGC CTGAGGA	CAGAGGGCCATC TCAGGTT	CACCATTAGCCACCA GCAACTGCT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
PMP22.1	NM_000304	NM_000304.2	PMP22	6254	CCATCTACACGGTG AGGCA	TGTAGGCGAAAC CGTAGGA	AATCCGAGTTGAGAT GCCACTCCG
PPAP2B.1	NM_003713	NM_003713.3	PPAP2B	6457	ACAAGCACCATCCC AGTGA	CACGAAGAAAAC ATGCAGCAG	ACCAGGGCTCCTTG AGCAAATCCT
PPARG.3	NM_005037	NM_005037.3	PPARG	1086	TGACTTTATGGAGC CCAAGTT	GCCAAGTCGCTG TCATCTAA	TTCCAGTGCATTGAA CTTCACAGCA
PPP1R3C.1	NM_005398	NM_005398.4	PPP1R3C	6320	TTCCTTCCCTCTCA ATCCAC	CACAGCTTTCCAT CACCATC	CCTTCCTCAACTTTT CCTTGCCCA
PPP2CA.1	NM_002715	NM_002715.2	PPP2CA	3879	GCAATCATGGAAC TGACGA	ATGTGGCTCGCC TCTACG	TTTCTTGCAGTTTGA CCCAGCACC
PRCC.1	NM_005973	NM_005973.4	PRCC	6002	GAGGAAGAGGAGG CGGTG	CAGGGAGAGAAG CGAACAA	CTACATCTGGGCCC GCTTTAGGG
PRKCA.1	NM_002737	NM_002737.1	PRKCA	2626	CAAGCAATGCGTCA TCAATGT	GTAATCCGCCC CCTCTTCT	CAGCCTCTGCGGAA TGGATCACACT
PRKCB1.1	NM_002738	NM_002738.5	PRKCB	3739	GACCCAGCTCCACT CTG	CCCATTACGTAC TCCATCA	CCAGACCATGGACC GCCTGTACTT
PRKCD.2	NM_006254	NM_006254.1	PRKCD	626	CTGACACTTGCCGC AGAGAA	AGGTGGTCCTTG GTCTGGAA	CCCTTTCTCACCCAC CTCATCTGCAC
PRKCH.1	NM_006255	NM_006255.3	PRKCH	4370	CTCCACCTATGAGC GTCTGTC	CACACTTTCCCTC CTTTTGG	TCCTGTTAACATCCC AAGCCCACA
PRO2000.3	NM_014109	NM_014109.2	ATAD2	1666	ATTGGAAAAACCTC GTCACC	TCGGTATCTTGGT CTTGCAG	CCCAACATATTTTAT AGTGGCCCAGC
PROM1.1	NM_006017	NM_006017.1	PROM1	4516	CTATGACAGGCATG CCACC	CTCCAACCATGAG GAAGACG	ACCCGAGGCTGTGT CTCCAACAC
PROM2.1	NM_144707	NM_144707.1	PROM2	5108	CTTCAGCGCATCCA CTACC	CCATGCTGGTCTT CACCAC	CTTCCTCGTTCAGAT CCAGAGGCC
PRPS2.1	NM_001039091	NM_001039091.1	PRPS2	4694	CACTGCACCAAGAT TCAGGT	ATTGTGTGTCCTT CGGATTG	TTGACATTTCCATGA TCTTGGCCG
PRSS8.1	NM_002773	NM_002773.2	PRSS8	4742	GTACACTCTGGCCT CCAGCTA	CCACACGAGGCT GGAGTT	TCCTGGATCCAAAGC AAGGTGACA
PSMA7.1	NM_002792	NM_002792.2	PSMA7	6255	GCCAACTGCAGG ATGAAAG	AGGCCATGCAGA CGTTGT	CCAAAGCACAGATCT TCCGCACTG
PSMB8.1	NM_148919	NM_148919.3	PSMB8	6461	CAGTGGCTATCGG CCTAATC	TAAGCAATAGCCC TGCGG	CAAGGTCATAGGCC TCTTCAGGGC
PSMB9.1	NM_148954	NM_148954.2	PSMB9	6462	GGGGTGTCTATCTAC CTGGTC	CATTGCCCAAGAT GACTCG	TGGTCCACACCGGC AGCTGTAATA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
PTEN.2	NM_000314	NM_000314.1	PTEN	54	TGGCTAAGTGAAGA TGACAATCATG	TGCACATATCATT ACACCAGTTCGT	CCTTTCAGCTTTAC AGTGAATTGCTGCA
PTGIS.1	NM_000961	NM_000961.3	PTGIS	5429	CCACACTGGCATCT CCCT	GCCCATGGGATG AGAAACT	CCTTCTCCAGGGAC AGAAGCAGGA
PTHR1.1	NM_000316	NM_000316.1	PTH1R	2375	CGAGGTACAAGCT GAGATCAAGAA	GCGTGCCTTTTCG CTTGAA	CCAGTGCCAGTGTC CAGCGGCT
PTK2.1	NM_005607	NM_005607.3	PTK2	2678	GACCGGTGGAATG ATAAGGT	CTGGACATCTCGA TGACAGC	ACCAGGCCCGTCAC ATTCTCGTAC
PTK2B.1	NM_004103	NM_004103.3	PTK2B	2883	CAAGCCCAGCCGA CCTAAG	GAACCTGGAAC GCAGCTTTG	CTCCGCAAACCAACC TCCTGGCT
PTN.1	NM_002825	NM_002825.5	PTN	3964	CCTTCCAGTCCAAA AATCCC	CCCCTCTCTCCAC TTTGGAT	TTCTCTGCTCTGGG GCTCTCTTG
PTPNS1.1	NM_080792	NM_080792.1	SIRPA	2896	CTCCAGCTAGCACT AAGCAACATC	TTTCAAGATTGCA CGTTTCACAT	TCTCAGTAATTTACA GGCGTCCACAG
PTPRB.1	NM_002837	NM_002837.2	PTPRB	3881	GATATGCGGTGAG GAACAGC	CTGGCCACACCG TATAGTGA	ATGCACACAGACTCA TCCGCCACT
PTPRC.1	NM_080921	NM_080921.2	PTPRC	6450	TGGCCGTCAATGG AAGAG	GGACATCTTTTGT GCTGGTTG	CAACATCTCCAAAAG CCCGAGTGC
PTPRG.1	NM_002841	NM_002841.2	PTPRG	2682	GGACAGCGACAAA GACTTGA	GGACTCGGAACA GGTAAAGG	CACCATTAGCCATGT CTCACCCGA
PTTG1.2	NM_004219	NM_004219.2	PTTG1	1724	GGCTACTCTGATCT ATGTTGATAAGGAA	GCTTCAGCCCATC CTTAGCA	CACACGGGTGCCTG GTTCTCCA
PVALB.1	NM_002854	NM_002854.2	PVALB	4316	AAACCAAGATGCTG ATGGCT	CAGCCACCAGAG TGGAGAA	TTTTGCCGTCCCAT CTTTGTCTC
PXDN.1	NM_012293	NM_012293.1	PXDN	6257	GCTGCTCAAGCTGA ACCC	ACCCACGATCTTC CTGGTC	ACTGGGACGGCGAC ACCATCTACT
RAC1.3	NM_006908	NM_006908.3	RAC1	2698	TGTTGTAAATGTCT CAGCCCC	TTGAGCAAAGCGT ACAAAGG	CGTCTTGGTCCTGT CCCTTGGA
RAD51.1	NM_002875	NM_002875.2	RAD51	3976	AGACTACTCGGGTC GAGGTG	AGCATCCGCAGA AACCTG	CTTTCAGCCAGGCA GATGCACTTG
RAF1.3	NM_002880	NM_002880.1	RAF1	2130	CGTCGTATGCGAG AGTCTGT	TGAAGGCGTGAG GTGTAGAA	TCCAGGATGCCTGTT AGTTCTCAGCA
RALBP1.1	NM_006788	NM_006788.2	RALBP1	2105	GGTGTGAGATATAA ATGTGCAAATGC	TTCGATATTGCCA GCAGCTATAAA	TGCTGTCTGTCTGG TCTCAGTACGTTCA
RARB.2	NM_016152	NM_016152.2	RARB	687	TGCCTGGACATCCT GATTCT	AAGGCCGTCTGA GAAAGTCA	TGCACCAGGTATACC CCAGAACAAGA

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
RASSF1.1	NM_007182	NM_007182.4	RASSF1	6658	AGGGCACGTGAAG TCATTG	AAAGAGTGCAAAC TTGCGG	CACCACCAAGAACTT TCGCAGCAG
RB1.1	NM_000321	NM_000321.1	RB1	956	CGAAGCCCTTACAA GTTTCC	GGACTCTTCAGG GGTGAAAT	CCCTTACGGATTCTCCT GGAGGGAAC
RBM35A.1	NM_017697	NM_017697.2	ESRP1	5109	TGGTTTTGAATCAC CAGGG	CTCTGTCCGCAG ACTTCATCT	CGCCCATCAGGAGA TGCCTTTATC
REG4.1	NM_032044	NM_032044.2	REG4	3226	TGCTAACTCCTGCA CAGCC	TGCTAGGTTTCCC CTCTGAA	TCCTCTTCCTTTCTG CTAGCCTGGC
RET.1	NM_020630	NM_020630.3	RET	5001	GCCTGTGCAGTTCT TGTGC	GGAAGGGCAGAC CCTCAC	AACATCAGCGTGCC CTACAGGCTC
RGS1.1	NM_002922	NM_002922.3	RGS1	6258	TGCCCTGTAAAGCA GAAGAGAT	CTCGAGTGCGGA AGTCAATA	AGCAGCATCTGAATG CACAAATGCT
RGS5.1	NM_003617	NM_003617.1	RGS5	2004	TTCAAACGGAGGCT CCTAAAGAG	GAAGGTTCCACC AGGTTCTTCA	AATATTGACCACTTC ACTAAGGACATCACA
RHEB.2	NM_005614	NM_005614.3	RHEB	6609	GATGATTGAGAACA GCCTTGC	GCTCCCAAGACT CTGACACA	TGTCAGTGCCTAGA ACACCCTGGAGTT
RhoB.1	NM_004040	NM_004040.2	RHOB	2951	AAGCATGAACAGGA CTTGACC	CCTCCCAAGTCA GTTGC	CTTTCCAACCCCTGG GGAAGACAT
rhoC.1	NM_175744	NM_175744.1	RHOC	773	CCCGTTCCGGTCTGA GGAA	GAGCACTCAAGG TAGCCAAAGG	TCCGGTTCGCCATGT CCCG
RIPK1.1	NM_003804	NM_003804.3	RIPK1	6259	AGTACCTTCAAGCC GGTCAA	AAGTCCCTGGGA ACTGTGC	CAGCCACAGAACAG CCTGGTTCAC
RND3.1	NM_005168	NM_005168.3	RND3	5381	TCGGAATTGGACTT GGGAG	CTGGTTACTCCCC TCCAACA	TTTAAAGCCTGACTC CTCACCGCG
ROCK1.1	NM_005406	NM_005406.1	ROCK1	2959	TGTGCACATAGGAA TGAGCTTC	GTTTAGCACGCAA TTGCTCA	TCACTCTCTTTGCTG GCCAACTGC
ROCK2.1	NM_004850	NM_004850.3	ROCK2	2992	GATCCGAGACCCT CGCTC	AGGACCAAGGAA TTTAAGCCA	CCCATCAACGTGGA GAGCTTGCT
RPLP1.1	NM_213725	NM_213725.1	RPLP1	5478	CAAGGTGCTCGGT CCTTC	GTCGCCGGATGA AGTGAG	CCTCACCCCAACGC AGCCTTAGCT
RPS23.1	NM_001025	NM_001025.1	RPS23	3320	GTTCTGTTGCTGG ATTTGG	CCTTAAAGCGGA CTCCAGG	ATCACCAACAGCATG ACCTTTGCG
RPS27A.1	NM_002954	NM_002954.3	RPS27A	6329	CTTACGGGGAAGA CCATCAC	TCCTGGATCTTGG CCTTTAC	TCGTATCCGAGGGTT CAACCTCG
RPS6KAI.1	NM_002953	NM_002953.3	RPS6KA1	3865	GCTCATGGAGCTA GTGCCTC	CGGCTGAAGTCC AGCTTCT	ACCCGGAGAATGGA CAGACCTCAG

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RPS6KB1.3	NM_003161	NM_003161.1	RPS6KB1	928	GCTCATTATGAAAA ACATCCCAAAC	AAGAAACAGAAGT TGTCTGGCTTTCT	CACACCAACCAATAA TTTCGCATT
RRM1.2	NM_001033	NM_001033.1	RRM1	1000	GGGCTACTGGCAG CTACATT	CTCTCAGCATCG GTACAAGG	CATTGGAATTGCCAT TAGTCCCAGC
RRM2.1	NM_001034	NM_001034.1	RRM2	2546	CAGCGGGATTAAAC AGTCCT	ATCTGCGTTGAAG CAGTGAG	CCAGCACAGCCAGT TAAAGATGCA
RUNX1.1	NM_001754	NM_001754.3	RUNX1	6067	AACAGAGACATTGC CAACCA	GTGATTTGCCCA GGAAAGTTT	TTGGATCTGCTTGCT GTCCAAACC
S100A1.1	NM_006271	NM_006271.1	S100A1	2851	TGGACAAGGTGAT GAAGGAG	AGCACCATATACT CCTGGAA	CCTCCCCGTCTCCAT TCTCGTCTA
S100A10.1	NM_002966	NM_002966.1	S100A10	3579	ACACCAAAAATGCCA TCTCAA	TTTATCCCCAGCG AATTTGT	CACGCCATGGAAAC CATGATGTTT
S100A2.1	NM_005978	NM_005978.2	S100A2	2369	TGGCTGTGCTGGT CACTACCT	TCCCCCTTACTCA GCTTGAAC	CACAAGTACTCCTGC CAAGAGGGCGAC
SAA2.2	NM_030754	NM_030754.2	SAA2	6655	CTACAGCACAGATC AGCACCA	TGCTGACACTCAG GACCAAG	AGCTTCTCACGGGC CTGGTTTTCT
SCN4B.1	NM_174934	NM_174934.3	SCN4B	7223	GCCTTCCTGGAGTA CCCG	GTGGCCCAATTC CCCAAGT	TGCTCCCTATGCCTT TCCAAGCAT
SCNN1A.2	NM_001038	NM_001038.4	SCNN1A	3263	ATCAACATCCTGTC GAGGCT	GAAGTTGCCCAG CGTGTC	AGAGACTCTGCCATC CCTGGAGGA
SDHA.1	NM_004168	NM_004168.1	SDHA	5443	GCAGAACTGAAGAT GGGAAGAT	CCCTTTCCAACT TGAGGC	CTGTCCACCAAATGC ACGCTGATA
SDPR.1	NM_004657	NM_004657.4	SDPR	3877	ACCAGCACAGATG GAGCA	GGTCATTCTGGAT GCCCTT	CGGAGCCCTCCAAA CTGATCTGTC
SELE.1	NM_000450	NM_000450.1	SELE	5383	AACTGGTCTGGC CTGCTAC	AAAGTCCAGCTAC CAAGGGAA	CCTGTGAAGCTCCC ACTGAGTCCA
SELENBP1.1	NM_003944	NM_003944.2	SELENBP1	6200	GGTACCAGCCTCG ACACAA	CCATCTCGTAAGA CATTGGGA	TGCCCCACTCAGTGCT GATCATGAC
SELL.1	NM_000655	NM_000655.3	SELL	5483	TGCAACTGTGATGT GGGG	CCTCCAAAGGCT CACACTG	CACAACTGACACTG GGGCCATA
SELPLG.1	NM_003006	NM_003006.3	SELPLG	6316	TGGCCACTATCTTC TTCGTG	GTAATTACGCACG GGGTACA	CACTGTGGTGCTGG CGGTCC
SEMA3B.1	NM_004636	NM_004636.1	SEMA3B	1013	GCTCCAGGATGTGT TTCTGTTG	ACGTGGAGAAGA CGGCATAGA	TCGCGGGACCACCG GACC
SEMA3C.1	NM_006379	NM_006379.2	SEMA3C	5409	ATGGCCATTCTGT TCCAG	GTCTCACATCTTG TCTTCGGC	CCTCCGTTTCCCAGT TGGGTAGAA

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SEMA3F.3	NM_004186	NM_004186.1	SEMA3F	1008	CGCGAGCCCCTCA TTATACA	CACTCGCCGTTG ACATCCT	CTCCCCACAGCGCA TCGAGGAA
SEMA5B.1	NM_001031702	NM_001031702.2	SEMA5B	5003	CTCGAGGACAGCT CCAACAT	TCACATTCCGCAC AGGAC	AGCCTCTGGACCCA GAACATCACC
SERPINA5.1	NM_000624	NM_000624.3	SERPINA5	6201	CAGCATGGTAGTG GCAAAGA	CTTTGTGGCACTG AGCTGG	AGGTCCAGAGTCCT GGCCCTTGAT
SFN.1	NM_006142	NM_006142.3	SFN	3580	GAGAGAGCCAGTC TGATCCA	AGGCTGCCATGT CCTCATA	CTGCTCTGCCAGCTT GGCCTTC
SGK.1	NM_005627	NM_005627.2	SGK1	2960	TCCGCAAGACACCT CCTG	TGAAGTCATCCTT GGCCC	TGTCCTGTCTTCTG CAGGAGGC
SHANK3.1	XM_037493	XM_037493.5		3887	CTGTGCCCTCTACA ACCAGG	GGACATCCCTGTT AGCTCCA	AGCTGTGCTCGTGT CCTGCTCTTC
SHC1.1	NM_003029	NM_003029.3	SHC1	2342	CCAACACCTTCTTG GCTTCT	CTGTTATCCCAAC CCAAACC	CCTGTGTTCTTGCTG AGCACCCCTC
SILV.1	NM_006928	NM_006928.3	SILV	4113	CCGCATCTTCTGCT CTTGT	ACTCAGACCTGCT GCCCCA	TTGGTGAGAATAGCC CCCTCCTCA
SKIL.1	NM_005414	NM_005414.2	SKIL	5272	AGAGGCTGAATATG CAGGACA	CTATCGGCCTCA GCATGG	CCAATCTCTGCCTCA GTTCTGCCA
SLC13A3.1	NM_022829	NM_022829.4	SLC13A3	6202	CTTGCCCTCCAACA AGGTC	AGCCCACTGAGG AAGAGGA	CCCCAGTACTTCCT CGACACCAA
SLC16A3.1	NM_004207	NM_004207.1	SLC16A3	4569	ATGCGACCCACGT CTACAT	AATCAGGGAGGA GGTGAGC	CCCCGCCAGGATGA ACACGTAC
SLC22A3.1	NM_021977	NM_021977.2	SLC22A3	6506	ATCGTCAGCGAGTT TGACCT	CAGGATGGCTTG GGTGAG	CAGCATCCACGCATT GACACAGAC
SLC22A6.1	NM_153277	NM_153277.1	SLC22A6	6463	TCCGCCACCTCTTC CTCT	GACCAGCCCATA GTATGCAAAG	CTCTCCATGCTGTGG TTTGCCACT
SLC2A1.1	NM_006516	NM_006516.1	SLC2A1	2966	GCCTGAGTCTCCTG TGCC	AGTCTCCACCCTC AGGCAT	ACATCCCAGGCTTCA CCCTGAATG
SLC34A1.1	NM_003052	NM_003052.3	SLC34A1	6203	GCTGAGACCCACT GACCTG	AGCCTCTCTCCGT AGGACAA	TCCTGGGCACCCAC TATGAGGTCT
SLC7A5.2	NM_003486	NM_003486.4	SLC7A5	3268	GCGCAGAGGCCAG TTAAA	AGCTGAGCTGTG GGTTGC	AGATCACCTCCTCGA ACCCACTCC
SLC9A1.1	NM_003047	NM_003047.2	SLC9A1	5385	CTTCGAGATCTCCC TCTGGA	AGTGGGGATCAC ATGGAAAC	CCTTCTGGCCTGCCT CATGAAGAT
SLIT2.2	NM_004787	NM_004787.1	SLIT2	3708	TTTACCGATGCACC TGTCC	ATGCAGGCATGA ATTGGG	CACAGTCCTGCCCC TTGAAACCAT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
SNAI1.1	NM_005985	NM_005985.2	SNAI1	2205	CCCAATCGGAAGC CTAACTA	GTAGGGCTGCTG GAAGGTAA	TCTGGATTAGAGTCC TGCAGCTCGC
SNRK.1	NM_017719	NM_017719.4	SNRK	6710	GAGGAAAAGTCAG GGCCG	GCCGGCTTTCAG AATCATC	CCAGCTGCAGTAGTT CGGAGACCA
SOD1.1	NM_000454	NM_000454.3	SOD1	2742	TGAAGAGAGGCAT GTTGGAG	AATAGACACATCG GCCACAC	TTTGTGAGCAGTCAC ATTGCCCAA
SP3.1	NM_001017371	NM_001017371.3	SP3	5430	TCAAGAGTCTCAGC AGCCAA	CCATGGATTGTCT GTGGTGT	CAGTCAAGCCCCAAT TGTGCAAGG
SPARC.1	NM_003118	NM_003118.1	SPARC	2378	TCTTCCCTGTACAC TGGCAGTTC	AGCTCGGTGTGG GAGAGGTA	TGGACCAGCACCCC ATTGACGG
SPARCL1.1	NM_004684	NM_004684.2	SPARCL1	3904	GGCACAGTGCAAG TGATGA	GATTGAGCTCTCT CGGCCT	ACTTCATCCCAAGCC AGGCCTTTC
SPAST.1	NM_014946	NM_014946.3	SPAST	4033	CCTGAGTTGTTTAC AGGGC	ATTCCCAGGTGG ACCAAAG	TAACAGCCCTCTGG CAGGAGCTCT
SPHK1.1	NM_021972	NM_021972.2	SPHK1	4178	GGCAGCTTCCTTGA ACCAT	GCAGTTGGTCAG GAGGTCTT	TGGTGACCTGCTCAT AGCCAGCAT
SPRY1.1	AK026960	AK026960.1		1051	CAGACCAGTCCCT GGTCATAGG	CCTTCAAGTCATC CACAATCAGTT	CTGGGTCCGGATTG CCCTTTCAG
SQSTM1.1	NM_003900	NM_003900.3	SQSTM1	4662	GGACCCGTCTACA GGTGAAC	GGGTCCAGAGAG CTTGGC	CAGTCCCTACAGATG CCAGAATCCG
STAT1.3	NM_007315	NM_007315.1	STAT1	530	GGGCTCAGCTTTCA GAAGTG	ACATGTTTCAGCTG GTCCACA	TGGCAGTTTTTCTTCT GTCACCAAAA
STAT3.1	NM_003150	NM_003150.1	STAT3	537	TCACATGCCACTTT GGTGTT	CTTGCAAGGAAGC GGCTATAC	TCCTGGGAGAGATT GACCAGCA
STAT5A.1	NM_003152	NM_003152.1	STAT5A	403	GAGGCGCTCAACA TGAAATTC	GCCAGGAACACG AGGTTCTC	CGGTTGCTCTGCACT TCGGCCT
STAT5B.2	NM_012448	NM_012448.1	STAT5B	857	CCAGTGGTGGTGA TCGTTCA	GCAAAAGCATTGT CCCAGAGA	CAGCCAGGACAACA ATGCGACGG
STC2.1	NM_003714	NM_003714.2	STC2	6507	AAGGAGGCCATCA CCCAC	AGATGGAGCACA GGCTTCC	TTCTGCTCACACTGA ACCTGCACG
STK11.1	NM_000455	NM_000455.3	STK11	3383	GGACTCGGAGACG CTGTG	GGGATCCTTCGC AACTTCTT	TTCTTGAGGATCTTG ACGGCCCTC
STK15.2	NM_003600	NM_003600.1	AURKA	341	CATCTTCCAGGAGG ACCACT	TCCGACCTTCAAT CATTTCA	CTCTGTGGCACCCT GGACTACCTG
STK4.1	NM_006282	NM_006282.2	STK4	5424	GAGCCATCTTCCTG CAACTT	CTGAGGTGCAAC CCAGTCA	ACCTCTTTCCTCAG ATGGGGAGC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
STMY3.3	NM_005940	NM_005940.2	MMP11	741	CCTGGAGGCTGCA ACATACC	TACAATGGCTTTG GAGGATAGCA	ATCCTCCTGAAGCCC TTTTCGCAGC
SUCLG1.1	NM_003849	NM_003849.2	SUCLG1	6205	CCAAGCCTGTAGTG TCCTTCA	CGGCATGACCCA TTCTTC	CCCAGGAGGAGCAG TTAAACCAGC
SULT1C2.1	NM_001056	NM_001056.3	SULT1C2	6206	GGGACCCAAAGCA TGAAAT	AGCACTGTTTCAT CCACCTTC	TTCCCATGAACTGCA TCACCTTCC
SURV.2	NM_001168	NM_001168.1	BIRC5	81	TGTTTTGATTCCCG GGCTTA	CAAAGCTGTCAG CTCTAGCAAAAG	TGCCTTCTTCCTCCC TCACTTCTCACCT
TACSTD2.1	NM_002353	NM_002353.2	TACSTD2	6335	ATCACCAACCGGAG AAAGTC	AAGCTCGGTTCT TTCTCAA	CCCCAGTTCCTTGA TCTCCACC
TAGLN.1	NM_003186	NM_003186.3	TAGLN	6073	GATGGAGCAGGTG GCTCAGT	AGTCTGGAACATG TCAGTCTTGATG	CCCATAGTCCTCAGC CGCCTTCAG
TAP1.1	NM_000593	NM_000593.5	TAP1	3966	GTATGCTGCTGAAA GTGGGAA	TCCCACTGCTTAC AGCCC	CACCAGCTGCCCCAC CAATGTAGAG
TCF4.1	NM_003199	NM_003199.1	TCF4	4097	CACACCCTGGAATG GGAG	ATGTGGCAACTTG GACCCT	CGCATCGAATCACAT GGGACAGAT
TCOF1.2	NM_001008656	NM_001008656.1	TCOF1	6719	AGCGAGGATGAGG ACGTG	CACCACATTGGTT CTGATGC	TCCCCGCTACACAGT GCTTGACTC
TEK.1	NM_000459	NM_000459.1	TEK	2345	ACTTCGGTGCTACT TAACAACCTTACATC	CCTGGGCCTTGG TGTTGAC	AGCTCGGACCACGT ACTGCTCCCTG
TERT.1	NM_003219	NM_003219.1		992	GACATGGAGAACAA GCTGTTTGC	GAGGTGTCACCA ACAAGAAATCAT	ACCAAACGCAGGAG CAGCCCG
TFAP2B.1	NM_003221	NM_003221.3	TFAP2B	6207	CGTGTGACGTGCG AGAGA	CCACACGCTCTCA GGACC	ATGGACGCGCCTTG CTCTTACTGT
TFAP2C.1	NM_003222	NM_003222.3	TFAP2C	4663	CATGCCTCACCAGA TGGA	CTGTCTGATCGTG CAGCAAC	CTGGTCGTCGACATT CTGCACCTC
TFPI.1	NM_006287	NM_006287.4	TFPI	6270	CCGAATGGTTTCCA GGTG	TTGCGGAGTCAG GGAGTTA	ATGGAACCCAGCTCA ATGCTGTGA
TGFA.2	NM_003236	NM_003236.1	TGFA	161	GGTGTGCCACAGA CCTTCCT	ACGGAGTTCTTGA CAGAGTTTTGA	TTGGCCTGTAATCAC CTGTGCAGCCTT
TGFb1.1	NM_000660	NM_000660.3	TGFB1	4041	CTGTATTTAAGGAC ACCCGTGC	TGACACAGAGATC CGCAGTC	TCTCTCCATCTTTAA TGGGGCCCC
TGFB2.2	NM_003238	NM_003238.1	TGFB2	2017	ACCAGTCCCCCAGA AGACTA	CCTGGTGCTGTT GTAGATGG	TCCTGAGCCCGAGG AAGTCCC
TGFBI.1	NM_000358	NM_000358.1	TGFBI	3768	GCTACGAGTGCTGT CCTGG	AGTGGTAGGGCT GCTGGAC	CCTTCTCCCCAGGG ACCTTTTCAT

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
TGFBR1.1	NM_004612	NM_004612.1	TGFBR1	3385	GTCATCACCTGGCC TTGG	GCAGACGAAGCA CACTGGT	AGCAATGACAGCTG CCAGTTCCAC
TGFBR2.3	NM_003242	NM_003242.2	TGFBR2	864	AACACCAATGGGT CCATCT	CCTCTTCATCAGG CCAAACT	TTCTGGGCTCCTGAT TGCTCAAGC
THBD.1	NM_000361	NM_000361.2	THBD	4050	AGATCTGCGACGG ACTGC	GGAAATGACATC GGCAGC	CACCTAATGACAGTG CGCTCCTCG
THBS1.1	NM_003246	NM_003246.1	THBS1	2348	CATCCGCAAAGTGA CTGAAGAG	GTACTGAACTCCG TTGTGATAGCATA G	CCAATGAGCTGAGG CGGCCTCC
TIMP1.1	NM_003254	NM_003254.2	TIMP1	6075	TCCCTGCGGTCCC AGATAG	GTGGGAACAGGG TGGACACT	ATCCTGCCCGGAGT GGAAGCTGAAGC
TIMP2.1	NM_003255	NM_003255.2	TIMP2	606	TCACCCTCTGTGAC TTCATCGT	TGTGGTTCAGGC TCTTCTTCTG	CCCTGGGACACCCT GAGCACCA
TIMP3.3	NM_000362	NM_000362.2	TIMP3	593	CTACCTGCCTTGCT TTGTGA	ACCGAAATTGGA GAGCATGT	CCAAGAACGAGTGT CTCTGGACCG
TK1.2	NM_003258	NM_003258.1	TK1	264	GCCGGGAAGACCG TAATTGT	CAGCGGCACCAG GTTCAG	CAAATGGCTTCCTCT GGAAGGTCCCA
TLR3.1	NM_003265	NM_003265.2	TLR3	6289	GGTTGGGCCACCT AGAAGT	CCATTCTGGCCT GTGAG	CTTGCCCAATTTTCT TAAGGCCCA
TMEM27.1	NM_020665	NM_020665.2	TMEM27	3878	CCCTGAAAGAATGT TGTGGC	TCTGCACCTGGTT GACAGAG	TCTGGTGACTGCCAT TCATGCTGA
TMEM47.1	NM_031442	NM_031442.3	TMEM47	6713	GGATTCCACTGTTA GAGCCCTT	GCAAATAACCAAC AGCCAATG	CCGCCTGCTTATCCT ACCCAATGA
TMSB10.1	NM_021103	NM_021103.3	TMSB10	6076	GAAATCGCCAGCTT CGATAA	GTCGGCAGGGTG TTCTTCT	CGTCTCCGTTTTTCTT CAGCTTGGC
TNF.1	NM_000594	NM_000594.1	TNF	2852	GGAGAAGGGTGAC CGACTCA	TGCCCAGACTCG GCAAAG	CGCTGAGATCAATC GGCCCGACTA
TNFAIP3.1	NM_006290	NM_006290.2	TNFAIP3	6290	ATCGTCTTGGCTGA GAAAGG	GTGGAATGGCTC TGGCTTC	CAACCCACGCGACTT GTGTGTCTT
TNFAIP6.1	NM_007115	NM_007115.2	TNFAIP6	6291	AGGAGTGAAAGAT GGGATGC	CTGTAAAGACGC CACCACAC	ATTGCTACAACCCAC ACGCAAAGG
TNFRSF10C.3	NM_003841	NM_003841.2	TNFRSF10C	6652	GGAGTTTGACCAGA GATGCAA	CTCTGTCCCCAGA GTTCCC	AACGGTAGGAAGCG CTCCTTCACC
TNFRSF10D.1	NM_003840	NM_003840.3	TNFRSF10D	4406	CCTCTCGCTTCTGG TGGTC	GCTCAGGAATCTC TGCCCTA	AGGCATCCCAGGGA CTCAGTTCAC
TNFRSF11B.1	NM_002546	NM_002546.2	TNFRSF11B	4675	TGGCGACCAAGAC ACCTT	GGGAAAGTGGTA CGTCTTTGAG	AGGGCCTAATGCAC GCACTAAAGC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
TNFRSF1A.1	NM_001065	NM_001065.2	TNFRSF1A	4943	ACTGCCCTGAGCC CAAAT	GTCAGGCACGGT GGAGAG	TGCCAGACAGCTAT GGCCTCTCAC
TNFSF12.1	NM_003809	NM_003809.2	TNFSF12	2987	TAGGCCAGGAGTT CCCAA	CACAGGGAATTCT CAAGGGA	TTGTCTTGTTTCTCG CCCCTCACA
TNFSF13B.1	NM_006573	NM_006573.3	TNFSF13B	4944	CCTACGCCATGGG ACATC	TCGAAACAAAGTC ACCAGACTC	TCCCCAAAGACATGG ACCTTCTTCC
TNFSF7.1	NM_001252	NM_001252.2	CD70	4101	CCAACCTCACTGG GACACTT	ACCCACTGCACTC CAAAGAA	TGCCTTCCCGAAACA CTGATGAGA
TNIP2.1	NM_024309	NM_024309.2	TNIP2	3872	CATGTCAGAAAGG GCCGA	GCGACCTTTTCCT CCAGTT	AATCCTACTTTGAGC CCGTTCCCG
TOP2A.4	NM_001067	NM_001067.1	TOP2A	74	AATCCAAGGGGGA GAGTGAT	GTACAGATTTTGC CCGAGGA	CATATGGACTTTGAC TCAGCTGTGGC
TOP2B.2	NM_001068	NM_001068.1	TOP2B	75	TGTGGACATCTTCC CCTCAGA	CTAGCCCGACCG GTTCGT	TTCCCTACTGAGCCA CCTTCTCTG
TP.3	NM_001953	NM_001953.2	TYMP	91	CTATATGCAGCCAG AGATGTGACA	CCACGAGTTTCTT ACTGAGAATGG	ACAGCCTGCCACTCA TCACAGCC
TRAIL.1	NM_003810	NM_003810.1	TNFSF10	898	CTTCACAGTGCTCC TGCAGTCT	CATCTGCTTCAGC TCGTTGGT	AAGTACACGTAAGTT ACAGCCACACA
TS.1	NM_001071	NM_001071.1	TYMS	76	GCCTCGGTGTGCC TTTCA	CGTGATGTGCGC AATCATG	CATCGCCAGCTACG CCCTGCTC
TSC1.1	NM_000368	NM_000368.3	TSC1	6292	TCACCAAATCTCAG CCCG	GTGTCAGCATAAG GGCTGGT	TTTCCTCATCGTTCA GCCGATGTC
TSC2.1	NM_000548	NM_000548	TSC2	5132	CACAGTGGCCTCTT TCTCCT	CAGGAAACGCTC CTGTGC	TACCAGTCCAGCTG CCAAGGACAG
TSPAN7.2	NM_004615	NM_004615.3	TSPAN7	6721	ATCACTGGGGTGAT CCTGC	GGGAGATATAGG TGCCCAGAG	AAGTTTGCCCCAGAC TCCAACAGC
TSPAN8.1	NM_004616	NM_004616.2	TSPAN8	6317	CAGAAATCTCTGCA GGCAAGT	AATCCAGATGCC GTGAATTT	TGCTCCAGAGCATAT TGCAGGACA
TUBB.1	NM_001069	NM_001069.1	TUBB2A	2094	CGAGGACGAGGCT TAAAAAC	ACCATGCTTGAG GACAACAG	TCTCAGATCAATCGT GCATCCTTAGTGAA
TUSC2.1	NM_007275	NM_007275.1	TUSC2	6208	CACCAAGAACGGG CAGAA	CGATGCCCTGAG GAATCA	TCTTATGCACTCGCC TCAGCTTGG
tusc4.2	NM_006545	NM_006545.4	TUSC4	3764	GGAGGAGCTAAAT GCCTCAG	CCTTCAAGTGGAT GGTGTTG	ACTCATCAATGGGCA GAGTGACC
TXLNA.1	NM_175852	NM_175852.3	TXLNA	6209	GCCAGAACGGCTC AGTCT	ATGTCTTCCAGTT GGCGG	TCCTCAGAGACATCA CGAAGGGCC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
UBB.1	NM_018955	NM_018955.1	UBB	3303	GAGTCGACCCTGC ACCTG	GCGAATGCCATG ACTGAA	AATTAACAGCCACCC CTCAGGCG
UBE1C.1	NM_003968	NM_003968.3	UBA3	2575	GAATGCACGCTGG AACTTTA	CTGGTAGCCTGG GCATAGA	AATTTTCCCATGTGC ACCATTGCA
UBE2C.1	NM_007019	NM_007019.2	UBE2C	2550	TGTCTGGCGATAAA GGGATT	ATGGTCCCTACCC ATTTGAA	TCTGCCTTCCCTGAA TCAGACAACC
UBE2T.1	NM_014176	NM_014176.1	UBE2T	3882	TGTTCTCAAATTGC CACCAA	AGAGGTCAACACA GTTGCGA	AGGTGCTTGGAGAC CATCCCTCAA
UGCG.1	NM_003358	NM_003358.1	UGCG	6210	GGCAACTGACAAAC AGCCTT	AGGATCTACCCCT TTCAGTGG	CAAGCTCCCAGGTG TCTCTCTTCTGA
UMOD.1	NM_003361	NM_003361.2	UMOD	6211	GCGTGGACCTGGA TGAGT	TTACGCAGCTGCT GTTGG	CCATTCTGGAGCTC ACAACTGCT
upa.3	NM_002658	NM_002658.1	PLAU	89	GTGGATGTGCCCT GAAGGA	CTGCGGATCCAG GGTAAGAA	AAGCCAGGCGTCTA CACGAGAGTCTCAC
USP34.1	NM_014709	NM_014709.2	USP34	4040	AAGCTGTGATGGC CAAGC	GGAATGGCCACA ACTGAGA	TCCCAGGACCCTGA GGTTGCTTTA
VCAM1.1	NM_001078	NM_001078.2	VCAM1	1220	TGGCTTCAGGAGCT GAATACC	TGCTGTCGTGAT GAGAAAATAGTG	CAGGCACACACAGG TGGGACACAAAT
VCAN.1	NM_004385	NM_004385.2	VCAN	5979	CCTGCTACACAGCC ACAAG	AGAAAGCGCCTG AGGTCC	CCCACTGTGGAAGA CAAAGAGGCC
VDR.2	NM_000376	NM_000376.1	VDR	971	GCCCTGGATTTTCA AAAGAG	AGTTACAAGCCAG GGAAGGA	CAAGTCTGGATCTG GGACCCTTTCC
VEGF.1	NM_003376	NM_003376.3	VEGFA	7	CTGCTGTCTTGGGT GCATTG	GCAGCCTGGGAC CACTTG	TTGCCTTGCTGCTCT ACCTCCACCA
VEGFB.1	NM_003377	NM_003377.2	VEGFB	964	TGACGATGGCCTG GAGTGT	GGTACCGGATCA TGAGGATCTG	CTGGGCAGCACCAA GTCCGGA
VHL.1	NM_000551	NM_000551.2	VHL	4102	CGGTGTTGACTT GTCTGC	AAGACTTGTCCCT GCCTCAC	ATGCCTCAGTCTTCC CAAAGCAGG
VIM.3	NM_003380	NM_003380.1	VIM	339	TGCCCTTAAAGGAA CCAATGA	GCTTCAACGGCA AAGTTCTCTT	ATTTACGCATCTGG CGTTCCA
VTCN1.1	NM_024626	NM_024626.2	VTCN1	4754	ACAGTGGTCTGGG CATCC	GCTCAAAGCTGG TATTGGAGAC	CGAGAAGTTGGCTC CCTGGTCAAC
VTN.1	NM_000638	NM_000638.2	VTN	4502	AGTCAATCTTCGCA CACGG	GTAAGTGGAGCGAT GGAGCGT	TGGACACTGTGGAC CCTCCCTACC
VWF.1	NM_000552	NM_000552.3	VWF	6212	TGAAGCACAGTGC CCTCTC	CCAGTCTCCATT CACCGT	CTCCATGTCACTGTG CAGCTCGAC

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Gene	Accession Num	Sequence_ID	Official Symbol	Gene Version ID	F Primer Seq	R Primer Seq	Probe Seq
WIF.1	NM_007191	NM_007191.3	WIF1	6077	AACAAGCTGAGTGC CCAGG	CACTCGCAGATG CGTCTTT	TACAAAAGCCTCCAT TTCGGCACC
WISP1.1	NM_003882	NM_003882.2	WISP1	603	AGAGGCATCCATGA ACTTCACA	CAAACCTCCACAGT ACTTGGGTTGA	CGGGCTGCATCAGC ACACGC
WT1.1	NM_000378	NM_000378.3	WT1	6458	TGTACGGTCGGCAT CTGAG	TTATTGCAGCCTG GGTAAGC	CAGTGAGAAACGCC CCTTCATGTG
WWOX.5	NM_016373	NM_016373.1	WWOX	974	ATCGCAGCTGGTG GGTGAC	AGCTCCCTGTTG CATGGACTT	CTGCTGTTTACCTTG GCGAGGCCTTTC
XDH.1	NM_000379	NM_000379.3	XDH	5089	TGGTGGCAGACAT CCCTT	GCCACAACCTGTC CCAGTCTT	TGAAGCCAACCTTGT ATCTGGCCA
XIAP.1	NM_001167	NM_001167.1	XIAP	80	GCAGTTGGAAGAC ACAGGAAAGT	TGCGTGCGCACTA TTTTCAAGA	TCCCCAAATTGCAGA TTTATCAACGGC
XPNPEP2.2	NM_003399	NM_003399.5	XPNPEP2	6503	CACCCTGCACTGAA CATACC	AAGGAGGATGAA TGCAAAGG	CCTGCTGGCCCATT GCCTAGAA
YB-1.2	NM_004559	NM_004559.1	YBX1	395	AGACTGTGGAGTTT GATGTTGTTGA	GGAACACCACCA GGACCTGTAA	TTGCTGCCTCCGCA CCCTTTTCT
ZHX2.1	NM_014943	NM_014943.3	ZHX2	6215	GAGTACGACCAGTT AGCGGC	TCTCCTTGAACCA ACGCAC	ATCTCAGTTCGGACC AGGCCAGTC

Table B

Gene	Target Sequence Length	Amplicon Sequence
A-Catenin.2	78	CGTTCCGATCCTCTATACTGCATCCCAGGCATGCCTACAGCACCTGATGTGCGCAGCCTATAAGGCCAA
A2M.1	66	CAGGGACCT CTCTCCCGCCTTCCTAGCTGTCCCAGTGGAGAAGGAACAAGCGCCTCACTGCATCTGTGCAAACGG
AAMP.1	66	GTGTGGCAGGTGGACACTAAGGAGGAGGTCTGGTCCTTTGAAGCGGGAGACCTGGAGTGGATGGAG
ABCB1.5	77	AAACACCACTGGAGCATTGACTACCAGGCTCGCCAATGATGCTGCTCAAGTTAAAGGGGCTATAGGTTT
ACADSB.1	68	CAGGCTTG TGGCGGAGAACTAGCCATCAGCCTCCTGAAGCCTGCCATCATTGTTAATTTGAGGACTGGGCTGTCTT
ACE.1	67	CCGCTGTACGAGGATTTCACTGCCCTCAGCAATGAAGCCTACAAGCAGGACGGCTTCACAGACACGG
ACE2.1	66	TACAATGAGAGGCTCTGGGCTTGGGAAAGCTGGAGATCTGAGGTCGGCAAGCAGCTGAGGCCATTA
ADAM17.1	73	GAAGTGCCAGGAGGCGATTAATGCTACTTGCAAAGGCGTGTCTACTGCACAGGTAATAGCAGTGAGTG
ADAM8.1	67	CCCC GTCCTGTGTCCAGCCCACCCTTCCCAGTTCCTGTCTACACCCGGCAGGCACCAAAGCAGGTCATCA
ADAMTS1.1	73	GGACAGGTGCAAGCTCATCTGCCAAGCCAAAGGCATTGGCTACTTCTTCGTTTTGCAGCCCAAGGTTGT
ADAMTS2.1	66	AGAT GAGAATGTCTGCCGCTGGGCCTACCTCCAGCAGAAGCCAGACACGGGCCACGATGAATACCACGAT
ADAMTS4.1	71	TTTGACAAGTGCATGGTGTGCGGAGGGGACGGTTCTGGTTGCAGCAAGCAGTCAGGCTCCTTCAGGAA
ADAMTS5.1	79	ATT CACTGTGGCTCACGAAATCGGACATTTACTTGGCCTCTCCCATGACGATTCCAAATTCTGTGAAGAGACC
ADAMTS8.1	72	TTTGGTTCC GCGAGTTCAAAGTGTTGAGGCCAAGGTGATTGATGGCACCTGTGTGGGCCAGAAACACTGGCCATCT
ADAMTS9.1	66	GTG GCACAGGTTACACAACCCAACAGAATGTCCCTATAACGGGAGCCGGCGCGATGACTGCCAATGTGCG
ADD1.1	74	GTCTACCCAGCAGCTCCGCAAGGAGGGATGGCTGCCTTAAACATGAGTCTTGGTATGGTGACTCCTGTG
ADFP.1	67	AACGA AAGACCATCACCTCCGTGGCCATGACCAGTGCTCTGCCCATCATCCAGAAGCTAGAGCCGCAAATTG
ADH1B.1	84	AAGCCAACAAACCTTCCTTCTTAACCATTTCTACTGTGTACCTTTGCCATTGAGGAAAAATATTCCTGTGA
ADH6.1	68	CTTCTTGCATTTT TGTTGGGGAGTAAACACTTGGACCTCTTGATATCCACCATCTTGGGCCATGAAGGGGCTGGAATCGTT
ADM.1	75	TAAGCCACAAGCACACGGGGCTCCAGCCCCCCCCGAGTGGAAGTGCTCCCCACTTTCTTTAGGATTTAGG
AGR2.1	70	CGCCCA AGCCAACATGTGACTAATTGGAAGAAGAGCAAAGGGTGGTGACGTGTTGATGAGGCAGATGGAGATCAG

Table B

Gene	Target Sequence Length	Amplicon Sequence
AGT.1	73	GATCCAGCCTCACTATGCCTCTGACCTGGACAAGGTGGAGGGTCTCACTTTCCAGCAAACTCCCTCAA
AGTR1.1	67	CTGG AGCATTGATCGATACCTGGCTATTGTTACCCCAATGAAGTCCCGCCTTCGACGCACAATGCTTGTA
AHR.1	69	GCGGCATAGAGACCGACTTAATACAGAGTTGGACCGTTTGGCTAGCCTGCTGCCTTTCCACAAGATGT
AIF1.1	71	GACGTTTACGCTACCCTGACTTTCTCAGGATGATGCTGGGCAAGAGATCTGCCATCCTAAAAATGATCCTG
AKT1.3	71	A CGCTTCTATGGCGCTGAGATTGTGTCAGCCCTGGACTACCTGCACTCGGAGAAGAACGTGGTGTACCG
AKT2.3	71	GGA TCCTGCCACCCTTCAAACCTCAGGTCACGTCCGAGGTCGACACAAGGTACTTCGATGATGAATTTACCG
AKT3.2	75	CC TTGTCTCTGCCTTGGACTATCTACATTCCGGAAAGATTGTGTACCGTGATCTCAAGTTGGAGAATCTAAT
ALDH4.2	68	GCTGG GGACAGGGTAAGACCGTGATCCAAGCGGAGATTGACGCTGCAGCGGAACATCGACTTCTTCCGGTT
ALDH6A1.1	66	GGCTCTTCAACAGCAGTCCTTGTGGGAGAAGCCAAGAAGTGGCTGCCAGAGCTGGTGGAGCATGC
ALDOA.1	69	GCCTGTACGTGCCAGCTCCCCGACTGCCAGAGCCTCAACTGTCTCTGCTTCGAGATCAAGCTCCGATGA
ALDOB.1	80	CCCTCTACCAGAAGGACAGCCAGGGAAAGCTGTTTCAAGGAAAAGGGGATCGTGGTGG GAATCAAGTTA
ALOX12.1	67	AGTTCCTCAATGGTGCCAACCCCATGCTGTTGAGACGCTCGACCTCTCTGCCCTCCAGGCTAGTGCT
ALOX5.1	66	GAGCTGCAGGACTTCGTGAACGATGTCTACGTGTACGGCATGCGGGGCCGCAAGTCCTCAGGCTTC
AMACR1.1	71	GGACAGTCAGTTTTAGGGTTGCCTGTATCCAGTAACTCGGGGCCTGTTTCCCCGTGGGTCTCTGGGCTG
ANGPT1.1	71	TC TCTACTTGGGGTGACAGTGCTCACGTGGCTCGACTATAGAAAACCTCCACTGACTGTGCGGGCTTTAAAAAG
ANGPT2.1	69	G CCGTGAAAGCTGCTCTGTAAAAGCTGACACAGCCCTCCCAAGTGAGCAGGACTGTTCTTCCCACTGCAA
ANGPTL2.1	66	GCCATCTGCGTCAACTCCAAGGAGCCTGAGGTGCTTCTGGAGAACCAGTGCTAAGCAGGAGCTA
ANGPTL3.3	78	GTTGCGATTACTGGCAATGTCCCCAATGCAATCCCGGAAAACAAAGATTTGGTGTTTTCTACTTGGGATC
ANGPTL4.1	66	ACAAAGCA ATGACCTCAGATGGAGGCTGGACAGTAATTCAGAGGCGCCACGATGGCTCAGTGGACTTCAACCGG
ANGPTL7.1	67	CTGCACAGACTCCAACCTCAATGGAGTGTAATACCGCCTGGGTGAGCACAATAAGCACCTGGATGGC

Table B

Gene	Target Sequence Length	Amplicon Sequence
ANTXR1.1	67	CTCCAGGTGTACCTCCAACCCTAGCCTTCTCCACAGCTGCCTACAACAGAGTCTCCAGCCTTCTC
ANXA1.2	67	GCCCTATCCTACCTTCAATCCATCCTCGGATGTCGCTGCCTTGCCATAAGGCCATAATGGTTAAAGG
ANXA2.2	71	CAAGACACTAAGGGCGACTACCAGAAAGCGCTGCTGTACCTGTGTGGTGGAGATGACTGAAGCCCGAC
ANXA4.1	67	TGGGAGGGATGAAGGAAATTATCTGGACGATGCTCTCGTGAGACAGGATGCCAGGACCTGTATGAG
ANXA5.1	67	GCTCAAGCCTGGAAGATGACGTGGTGGGGGACACTTCAGGGTACTACCAGCGGATGTTGGTGGTTCT
AP-1 (JUN official).2	81	GACTGCAAAGATGGAACGACCTTCTATGACGATGCCCTCAACGCCTCGTTCCTCCCGTCCGAGAGCGG
AP1M2.1	67	ACCTTATGGCTA
APAF1.2	66	ACAACGACCGCACCATCTCCTTCATCCCGCCTGATGGTGACTTTGAGCTCATGTCATACCGCCTCAG
APC.4	66	CACAAGGAAGAAGCTGGTGAATGCAATTCAGCAGAAGCTCTCCAAATTGAAAGGTGAACCAGGATG
APOC1.3	69	GGACAGCAGGAATGTGTTTCTCCATACAGGTCACGGGGAGCCCAATGGTTCAGAAACAAATCGAGTGGGT
APOE.1	70	CCAGCCTGATAAAGGTCTGCGGGCAGGACAGGACCTCCCAACCAAGCCCTCCAGCAAGGATTCAGAG
APOL1.1	75	TG
APOLD1.1	73	GCCTCAAGAGCTGGTTCGAGCCCCTGGTGGAAAGACATGCAGCGCCAGTGGGCCGGGCTGGTGGAGAA
AQP1.1	66	GGTGCAGG
AREG.2	82	CGGACCAAGAACTGTGACCACAGGGCAGGGCAGCCACCAGGAGAGATATGCCTGGCAGGGGCCAGGA
ARF1.1	64	CAAAAT
ARG99.1	67	GAGCAGCTGGAGTCTCGGGTTCAGCTCTGCACCAAGTCCAGTCGTGGCCACGACCTCAAGATCTCT
ARGHEF18.1	71	GCTTGCTGTATGACCCCTGGCCACAGCCTTCCCTCTGCATTGACCTGGAGGGGAGAGGTCAGCCTT
ARHA.1	73	TGTGAGTGAAATGCCTTCTAGTAGTGAACCGTCCTCGGGAGCCGACTATGACTACTCAGAAGAGTATGAT
ARHGDI.1	66	AACGAACCACAA
ARRB1.1	69	CAGTAGAGATCCCCGCAACTCGCTTGCTCCTTGGGTACCCCTGCATTCCATAGCCATGTGCTTGT
	67	GCATGGGCTACTGCATCCTTTTTGTGCACGGACTGAGCAAGCTCTGCACCTGGCTGAATCGATGTGG
	71	ACTCTGCTTCCCAAGGGCAACCGTTGCTGTTCACACGCTCAGCCTGTCTGGGGGAGCGGGCCTCTAGC
	73	TTC
	66	GGTCTCCGTGCGTTCTCTCATTAGTCCACGGTCTGGTCTTCAGCTACCCGCCTTCGTCTCCGAGTTTG
	66	CGAC
	69	TGGTCCCTAGAACAAAGAGGCTTAAACCGGGCTTTCACCCAACCTGCTCCCTCTGATCCTCCATCA
	69	TGCAGGAACGCCTCATCAAGAAGCTGGGCGAGCACGCTTACCCTTTCACCTTTGAGATCCCTCCAAACC

Table B

Gene	Target Sequence Length	Amplicon Sequence
ASS1.1	85	CCCCCAGATAAAGGTCATTGCTCCCTGGAGGATGCCTGAATTCTACAACCGGTTCAAGGGCCGCAATGA CCTGATGGAGTACGCA
ATP1A1.1	67	AGAACGCCTATTTGGAGCTGGGGGGCCTCGGAGAACGAGTCCTAGGTTTCTGCCACCTCTTTCTGCC
ATP5E.1	66	CCGCTTTCGCTACAGCATGGTGGCCTACTGGAGACAGGCTGGACTCAGCTACATCCGATACTCCCA
ATP6V1B1.1	67	AACCATGGGGAACGTCTGCCTCTTCCTGAAC TTGGCCAATGACCCACGATCGAGCGGATCATCACC
AXL.1	66	TTGCAGCCCTGTCTTCCTACCTATCCACCTCCATCCCAGACAGGTCCCTCCCCTTCTCTGTGCAG CCGAGGCCCTGACTTCTTCACCCGAGTGGCGCTCTTCCGAGACTGGATCGATGGTGTCTCAACAACCC
AZU1.1	74	GGGAC GGCTCTTGTCGCTACTGTCCTTCGGGCTGGTGACAGGGAAGACATCACTGAGCCTGCCATCTGTGCTCT
B-Catenin.3	80	TCGTCATCTGA
B2M.4	67	GGGATCGAGACATGTAAGCAGCATCATGGAGGTTTGAAGATGCCGCATTTGGATTGGATGAATTCCA GGGTCAGGGGCCTCGAGATCGGGCTTGGGCCAGAGCATGTTCCAGATCCCAGAGTTTGAGCCGAGTG
BAD.1	73	AGCAG
BAG1.2	81	CGTTGTCAGCACTTGGAATACAAGATGGTTGCCGGGTCATGTTAATTGGGAAAAAGAACAGTCCACAGG AAGAGGTTGAAC
BAG2.1	69	CTAGGGGGCAAAAAGCATGACTGCTTTTTCTGTCTGGCATGGAATCACGCAGTCACCTTGGGCATTTAG
Bak.2	66	CCATTCCCACCATTCTACCTGAGGCCAGGACGTCTGGGGTGTGGGGATTGGTGGGTCTATGTTCCC CCGCCGTGGACACAGACTCCCCCGAGAGGTCTTTTTCCGAGTGGCAGCTGACATGTTTTCTGACGGCA
Bax.1	70	A
BBC3.2	83	CCTGGAGGGTCCTGTACAATCTCATCATGGGACTCCTGCCCTTACCCAGGGGCCACAGAGCCCCCGAG ATGGAGCCCAATTAG
Bcl2.2	73	CAGATGGACCTAGTACCCACTGAGATTTCCACGCCGAAGGACAGCGATGGGAAAAATGCCCTTAAATCA TAGG
BCL2A1.1	79	CCAGCCTCCATGTATCATCATGTGTCATAACTCAGTCAAGCTCAGTGAGCATTCTCAGCACATTGCCTCA ACAGCTTCA
BCL2L12.1	73	AACCCACCCCTGTCTTGGAGCTCCGGGTAGCTCTCAAACCTCGAGGCTGCGCACCCCTTTCCCGTCAGC TGAG
Bclx.2	70	CTTTTGTGGAACCTCTATGGGAACAATGCAGCAGCCGAGAGCCGAAAGGGGCCAGGAACGCTTCAACCGCT G
BCRP.1	74	TGTACTGGCGAAGAATATTTGGTAAAGCAGGGCATCGATCTCTCACCTGGGGCTTGTGGAAGAATCAC GTGGC
BFGF.3	77	CCAGGAAGAATGCTTAAGATGTGAGTGGATGGATCTCAATGACCTGGCGAAGACTGAAAATACAACCTCC CATCACCA

Table B

Gene	Target Sequence Length	Amplicon Sequence
BGN.1	66	GAGCTCCGCAAGGATGACTTCAAGGGTCTCCAGCACCTCTACGCCCTCGTCCTGGTGAACAACAAG
BHLHB3.1	68	AGGAAGATCCCTCGCAGCCAGGAAAGGAAGCTCCCTGAATCCTTGCGTCCCGAAGGACGGAGGTTCAA
BIK.1	70	C
BIN1.3	76	CCTGCAAAAAGGGAACAAGAGCCCTTCGCCTCCAGATGGCTCCCCTGCCGCCACCCCCGAGATCAGAGT CAACCACG
BLR1.1	67	GACCAAGCAGGAAGCTCAGACTGGTTGAGTTCAGGTAGCTGCCCCTGGCTCTGACCGAAACAGCGCT
BNIP3.1	68	CTGGACGGAGTAGCTCCAAGAGCTCTCACTGTGACAGCCCACCTCGCTCGCAGACACCACAAGATACC
BRCA1.1	65	TCAGGGGGGCTAGAAATCTGTTGCTATGGGCCCTTCACCAACATGCCACAGATCAACTGGAATGG
BTRC.1	63	GTTGGGACACAGTTGGTCTGCAGTCGGCCCAGGACGGTCTACTCAGCACAACCTGACTGCTTCA
BUB1.1	68	CCGAGGTTAATCCAGCACGTATGGGGCCAAGTGTAGGCTCCCAGCAGGAAGTGAAGAGCGCCATGTCTT
BUB3.1	73	CTGAAGCAGATGGTTCATCATTTCCTGGGCTGTAAACAAAGCGAGGTTAAGGTTAGACTCTTGGGAATC AGC
c-kit.2	75	GAGGCAACTGCTTATGGCTTAATTAAGTCAGATGCGGCCATGACTGTGCTGTAAAGATGCTCAAGCCG AGTGCC
C13orf15.1	84	TAGAATCTGCTGCCAGAGGGGACAAAGACGTGCACTCAACCTTCTACCAGGCCACTCTCAGGCTCACCT TAAAATCAGCCCTTG
C1QA.1	66	CGGTCATACCAACCAGGAAGAACCGTACCAGAACCACTCCGGCCGATTTCGTCTGCACTGTACCCG CCAGTGGCCTCACAGGACACCAGCTTCCCAGGAGGCGTCTGACACAGTATGATGATGAAGATCCCATGG
C1QB.1	70	G
C20 orf1.1	65	TCAGCTGTGAGCTGCGGATACCGCCCGGCAATGGGACCTGCTCTTAACCTCAAACCTAGGACCGT
C3.1	67	CGTGAAGGAGTGCAGAAAGAGGACATCCCACCTGCAGACCTCAGTGACCAAGTCCCGGACACCGAGT
C3AR1.1	66	AAGCCGCATCCCAGACTTGCTGAATCGGAATCTCTGGGGGTTGGGACCCAGCAAGGGCACTTAACA ATGTCTGAGTGTGAGGCGGGCGCTCTGAGATGCAGAGGGCAGAGCATCTCTGTCAACAGCATAAGGCC
C7.1	69	T
CA12.1	66	CTCTCTGAAGGTGTCCTGGCCAGCCCTGGAGAAGCACTGGTGTCTGCAGCACCCCTCAGTTCCTGT
CA2.1	69	CAACGTGGAGTTTGATGACTCTCAGGACAAAGCAGTGCTCAAGGGAGGACCCCTGGATGGCACTTACAG

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Gene	Target Sequence Length	Amplicon Sequence
CA9.3	72	ATCCTAGCCCTGGTTTTTGGCCTCCTTTTGCTGTCACCAGCGTCGCGTTCCTTGTGCAGATGAGAAGG CAG
CACNA2D1.1	68	CAAACATTAGCTGGGCCTGTTCCATGGCATAAACACTAAGGCCGAGACTCCTAAGGCACCCACTGGCTG CACTAAGGTTTTGAGACAGTTCCAGAAAGAACCCAAGCTCAAGACGCAGGACGAGCTCAGTTGTAGAGGG
CALD1.2	78	CTAATTCGC AGAAAGCCCACATAGAGAAGGATTTTATCGCTTTCTGCTCTTCCACACCAGATAATGTTTCTTGGAGACAT
CASP1.1	77	CCCCACA
CASP10.1	66	ACCTTTCTCTTGGCCGGATGTCCTCAGGGCTGGCAGATGCAGTAGACTGCAGTGGACAGTCCCCAC
CASP6.1	67	CCTCACACTGGTGAACAGGAAAGTTTCTCAGCGCCGAGTGGACTTTTGCAAAGACCCAAGTGCAATT
Caspase 3.1	66	TGAGCCTGAGCAGAGACATGACTCAGCCTGTTCCATGAAGGCAGAGCCATGGACCACGCAGGAAGG ATCCATTGATCTCACCAAGGTTTGGCCTCACAAGGACTACCCTCTCATCCCAGTTGGTAAACTGGTCTT
CAT.1	78	AAACCGGA GTGGCTCAACATTGTGTTCCATTTCAGCTGATCAGTGGGCCTCCAAGGAGGGGGCTGTAAAATGGAGGC
CAV1.1	74	CATTG
CAV2.1	66	CTTCCCTGGGACGACTTGCCAGCTCTGAGGCATGACAGTACGGGCCCCCAGAAGGGTGACCAGGAG
CCL18.1	68	GCTCCTGTGCACAAGTTGGTACCAACAAAGAGCTCTGCTGCCTCGTCTATA CCTCCTGGCAGATTCCA GAACGCATCATCCAGAGACTGCAGAGGACCTCAGCCAAGATGAAGCGCCGCAGCAGTTAACCTATGACC
CCL19.1	78	GTGCAGAGG
CCL20.1	69	CCATGTGCTGTACCAAGAGTTTGCTCCTGGCTGCTTTGATGTGCTGCTGCTACTCCACCTCTGCGGCG GGGTCCAGGAGTACGTGTATGACCTGGA ACTGAACTGAGCTGCTCAGAGACAGGAAGTCTTCAGGGAA
CCL4.2	70	GG
CCL5.2	65	AGGTTCTGAGCTCTGGCTTTGCCTTGGCTTTGCCAGGGCTCTGTGACCAGGAAGGAAGTCAGCAT TTCAGGTTGTTGCAGGAGACCATGTACATGACTGTCTCCATTATTGATCGGTT CATGCAGAATAATTGTG
CCNB1.2	84	TGCCCAAGAAGATG
CCND1.3	69	GCATGTTCGTGGCCTCTAAGATGAAGGAGACCATCCCCCTGACGGCCGAGAAGCTGTGCATCTACACCG AAAGAAGATGATGACCGGGTTTACCCAAACTCAACGTGCAAGCCTCGGATTATTGCACCATCCAGAGGC
CCNE1.1	71	TC
CCNE2 variant 1.1	85	GGTCACCAAGAAACATCAGTATGAAATTAGGAATTGTTGGCCACCTGTATTATCTGGGGGGATCAGTCCT TGCATTATCATTGAA

Table B

Gene	Target Sequence Length	Amplicon Sequence
CCNE2.2	82	ATGCTGTGGCTCCTTCCTAACTGGGGCTTTCTTGACATGTAGGTTGCTTGGTAATAACCTTTTTGTATATC
CCR1.1	66	ACAATTTGGGT TCCAAGACCCAATGGGAATTCACCTCACCACACCTGCAGCCTTCACTTTCCTCACGAAAGCCTACGA
CCR2.1	67	CTCGGGAATCCTGAAAACCCTGCTTCGGTGTGCGAAACGAGAAGAAGAGGCATAGGGCAGTGAGAGTC
CCR4.2	82	AGACCCTGGTGGAGCTAGAAGTCCTTCAGGACTGCACCTTTGAAAGATACTTGGACTATGCCATCCAGG CCACAGAAACTCT
CCR5.1	67	CAGACTGAATGGGGGTGGGGGGGGCGCCTTAGGTACTTATTCCAGATGCCTTCTCCAGACAAACCAG
CCR7.1	64	GGATGACATGCACTCAGCTCTTGGCTCCACTGGGATGGGAGGAGAGGACAAGGGAAATGTCAGG
CD105.1	75	GCAGGTGTCAGCAAGTATGATCAGCAATGAGGCGGTGGTCAATATCCTGTGCGAGCTCATCACCACAGCG GAAAAA
CD14.1	66	GTGTGCTAGCGTACTCCCGCCTCAAGGAACTGACGCTCGAGGACCTAAAGATAACCGGCACCATGC
CD18.2	81	CGTCAGGACCCACCATGTCTGCCCCATCACGCGGCCGAGACATGGCTTGGCCACAGCTCTTGAGGATG TCACCAATTAACC
CD1A.1	78	GGAGTGGAAGGAACTGGAACATTATTCCGTATACGCACCATTCGGTCATTTGAGGGAATTCGTAGATAC GCCCATGA
CD24.1	77	TCCAATAATGCCACCACCAAGGCGGCTGGTGGTGCCCTGCAGTCAACAGCCAGTCTCTTCGTGGTCTC ACTCTCTC
CD274.2	69	GCTGCATGATCAGCTATGGTGGTGCCGACTACAAGCGAATTACTGTGAAAGTCAATGCCCCATACAACA
CD31.3	75	TGTATTTCAAGACCTCTGTGCACTTATTTATGAACCTGCCCTGCTCCCACAGAACACAGCAATTCCTCAG GCTAA
CD34.1	67	CCACTGCACACACCTCAGAGGCTGTTCTTGGGGCCCTACACCTTGAGGAGGGGGCAGGTAAACTCCTG
CD36.1	67	GTAACCCAGGACGCTGAGGACAACACAGTCTCTTTCCTGCAGCCCAATGGTGCCATCTTCGAACCTT
CD3z.1	65	AGATGAAGTGGAAGGCGCTTTTCACCGCGGCCATCCTGCAGGCACAGTTGCCGATTACAGAGGCA
CD4.1	67	GTGCTGGAGTCGGGACTAACCCAGGTCCCTTGTCCCAAGTTCCACTGCTGCCTCTTGAATGCAGGGA
CD44.1	67	GGCACCCTGCTTATGAAGGAACTGGAACCCAGAAGCACACCCTCCCCTCATTACCATGAGCATC
CD44s.1	78	GACGAAGACAGTCCCTGGATCACCGACAGCACAGACAGAATCCCTGCTACCAGAGACCAAGACACATTC CACCCAGT
CD44v6.1	78	CTCATACCAGCCATCCAATGCAAGGAAGGACAACACCAAGCCCAGAGGACAGTTCCTGGACTGATTTCTT CAACCCAA

Table B

Gene	Target Sequence Length	Amplicon Sequence
CD53.1	72	CGACAGCATCCACCGTTACCACTCAGACAATAGCACCAAGGCAGCGTGGGACTCCATCCAGTCATTTCTGCA
CD68.2	74	TGGTTCCAGCCCTGTGTCCACCTCCAAGCCCAGATTGAGATTCGAGTCATGTACACAACCCAGGGTGGAGGAG
CD82.3	84	GTGCAGGCTCAGGTGAAGTGCTGCGGCTGGGTCAGCTTCTACAACCTGGACAGACAACGCTGAGCTCATGAATCGCCCTGAGGTC
CD8A.1	68	AGGGTGAGGTGCTTGAGTCTCCAACGGCAAGGGAACAAGTACTTCTTGATACCTGGGATACTGTGCCC GTTCCTCCGGTAGCTTTTCAGATGCTGACCTTGCGGATGGCGTTTCAGGTGGAGAAGGAAAAGGAGGCA
CD99.1	77	GTGATGGT
cdc25A.4	71	TCTTGCTGGCTACGCCTCTTCTGTCCCTGTTAGACGTCCTCCGTCCATATCAGAACTGTGCCACAATGCA G
CDC25B.1	85	AAACGAGCAGTTTGCCATCAGACGCTTCCAGTCTATGCCGGTGAGGCTGCTGGGCCACAGCCCCGTGCTTCGGAACATCACCAAC
CDH1.3	81	TGAGTGTCCCCCGGTATCTTCCCCGCCCTGCCAATCCCGATGAAATTGGAAATTTTATTGATGAAAATCT GAAAGCGGCTG
CDH13.1	67	GCTACTTCTCCACTGTCCCGTTCAGTCTGAATGCTGCCACAACCAGCCAGGCAGGTCCACAGAGAGG
CDH16.1	67	GACTGTCTGAATGGCCCAGGCAGCTCTAGCTGGGAGCTTGGCCTCTGGCTCCATCTGAGTCCCCTGG
CDH2.1	66	TGACCGATAAGGATCAACCCCATACACCAGCCTGGAACGCAGTGTACAGAATCAGTGGCGGAGATC
CDH5.1	67	ACAGGAGACGTGTTCCGCCATTGAGAGGCTGGACCGGGAGAATATCTCAGAGTACCACCTCACTGCTG
CDH6.1	66	ACACAGGCGACATACAGGCCACCAAGAGGCTGGACAGGGAAGAAAAACCCGTTTACATCCTTCGAG
CDK4.1	66	CCTTCCCATCAGCACAGTTCGTGAGGTGGCTTTACTGAGGCGACTGGAGGCTTTTGAGCATCCCAA
CDK6.1	67	AGTGCCCTGTCTCACCCTACTTCCAGGACCTGGAAAGGTGCAAAGAAAACCTGGATTCCCACCTGC AGCACTCACGCCCTAAGCGCACATTCATGTGGGCATTTCTTGCGAGCCTCGCAGCCTCCGGAAGCTGTC
CDKN2A.2	79	GACTTCATGA
CEACAM1.1	71	ACTTGCCGTGTTTCAGAGCACTCATTCTTCCCACCCCCAGTCCTGTCTATCACTCTAATTCGGATTTGCC A
CEBPA.1	66	TTGGTTTTTGCTCGGATACTTGCCAAAATGAGACTCTCCGTCCGCAGCTGGGGGAAGGGTCTGAGAC
CENPF.1	68	CTCCCGTCAACAGCGTTCTTTCCAAACACTGGACCAGGAGTGCATCCAGATGAAGGCCAGACTCACCC
CFLAR.1	66	GGACTTTTGTCCAGTGACAGCTGAGACAACAAGGACCACGGGAGGAGGTGTAGGAGAGAAGCGCCG

Table B

Gene	Target Sequence Length	Amplicon Sequence
CGA (CHGA official).3	76	CTGAAGGAGCTCCAAGACCTCGCTCTCCAAGGCGCCAAGGAGAGGGGCACATCAGCAGAAGAAACACAG CGGTTTTG
Chk1.2	82	GATAAATTGGTACAAGGGATCAGCTTTTCCCAGCCCACATGTCCTGATCATATGCTTTTGAATAGTCAGTT ACTTGGCACCC
Chk2.3	78	ATGTGGAACCCCCACCTACTTGGCGCCTGAAGTTCTTGTTTCTGTTGGGACTGCTGGGTATAACCGTGC TGTGGACTG
CIAP1.2	72	TGCCTGTGGTGGGAAGCTCAGTAACTGGGAACCAAAGGATGATGCTATGTCAGAACACCGGAGGCATTT TCC
cIAP2.2	86	GGATATTTCCGTGGCTCTTATTCAAACCTCTCCATCAAATCCTGTAAACTCCAGAGCAAATCAAGATTTTTTC TGCCTTGATGAGAAG
CLCNKB.1	67	GTGACCCTGAAGCTGTCCCCAGAGACTTCCCTGCATGAGGCACACAACCTCTTTGAGCTGTTGAACC
CLDN10.1	66	GGTCTGTGGATGAACTGCGCAGGTAACGCGTTGGGTTCTTTCCATTGCCGACCGCATTTTACTATC GGTCTGCCCTAGTCATCCTGGGAGGTGCACTGCTCTCCTGTTCTGCTGGGAATGAGAGCAAGGCTG
CLDN7.2	74	GGTAC
CLU.3	76	CCCCAGGATACCTACCACTACCTGCCCTTCAGCCTGCCCCACCGGAGGCCTCACTTCTTCTTTCCCAAG TCCCGCA
cMet.2	86	GACATTTCCAGTCTGTCAGTCAATGCCTCTCTGCCCCACCCTTTGTTCAGTGTGGCTGGTGCCACGACA AATGTGTGCGATCGGAG
cMYC.3	84	TCCCTCCACTCGGAAGGACTATCCTGCTGCCAAGAGGGTCAAGTTGGACAGTGTGAGAGTCCTGAGACA GATCAGCAACAACCG
COL18A1.1	67	AGCTGCCATCACGCCTACATCGTGCTCTGCATTGAGAACAGCTTCATGACTGCCTCCAAGTAGCCAC
COL1A1.1	68	GTGGCCATCCAGCTGACCTTCCTGCGCCTGATGTCCACCGAGGCCTCCCAGAACATCACCTACCACTG CAGCCAAGAAGTGGTATAGGAGCTCCAAGGACAAGAAACACGTCTGGCTAGGAGAACTATCAATGCTG
COL1A2.1	80	GCAGCCAGTTT
COL4A1.1	66	ACAAAGGCCTCCCAGGATTGGATGGCATCCCTGGTGTCAAAGGAGAAGCAGGTCTTCCTGGGACTC
COL4A2.1	67	CAACCCTGGTGATGTCTGCTACTATGCCAGCCGGAACGACAAGTCCTACTGGCTCTTACCACTGCG GGTCGAGGAACCCAAGGTCCGCCTGGTGCTACAGGATTTCTGGTCTGCGGGCAGAGTTGGACCTCC
COL5A2.2	72	AGGC
COL7A1.1	66	GGTGACAAAGGACCTCGGGGAGACAATGGGGACCCTGGTGACAAGGGCAGCAAGGGAGAGCCTGGT TCTGCAGAGTTGGAAGCACTCTATGGTGACATCGATGCTGTGGAGCTGTATCCTGCCCTTCTGGTAGAA
COX2.1	79	AAGCCTCGGC

Table B

Gene	Target Sequence Length	Amplicon Sequence
CP.1	73	CGTGAGTACACAGATGCCTCCTTCACAAATCGAAAGGAGAGAGGCCCTGAAGAAGAGCATCTTGGCATC
CPB2.1	67	CTGG GGCACATACGGATTCTTGCTGCCGGAGCGTTACATCAAACCCACCTGTAGAGAAGCTTTTGCCGCTG
CRADD.1	69	GATGGTGCCTCCAGCAACCGCTGGGGAGTGTGTCCCTGAGTCATGTGGGCTGAATCCTGACTTTCACTC
cripto (TDGF1 official).1	65	GGGTCTGTGCCCCATGACACCTGGCTGCCCAAGAAGTGTTCCCTGTGTAAATGCTGGCACGGTCA
CRP.1	66	GACGTGAACCACAGGGTGTCTGTGTCAGAGGAGCCCATCTCCCATCTCCCCAGCTCCCTATCTGGAG
		TGCAGCGGCTGATTGACAGTCAGATGGAGACCTCGTGCCAAATTACATTTGAGTTTGTAGACCAGGAAC
CSF1.1	74	AGTTG
		GAGCACAACCAAACCTACGAGTGCAGGGCCCCACAACAGCGTGGGGAGTGGCTCCTGGGCCTTCATACC
CSF1R.2	80	CATCTCTGCAGG
		GAACCTGAAGGACTTTCTGCTTGTCTATCCCCTTTGACTGCTGGGAGCCAGTCCAGGAGTGAGACCGGCC
CSF2.1	76	AGATGAG
CSF2RA.2	67	TACCACACCCAGCATTTCCTCCTGATCCCAGAGAAATCGGATCTGCGAACAGTGGCACCAGCCTCTAG
		CCCAGGCCTCTGTGTCTTCCCTGCATTTCTGAGTTTCATTCTCCTGCCTGTAGCAGTGAGAAAAAGCTC
CSF3.2	79	CTGTCCTCC
		GAGTTCAAGTGCCCTGACGGCGAGGTCATGAAGAAGAACATGATGTTTCATCAAGACCTGTGCCTGCCAT
CTGF.1	76	TACAACT
CTSB.1	62	GGCCGAGATCTACAAAAACGGCCCCCGTGGAGGGAGCTTTCTCTGTGTATTTCGGACTTCCTGC
		GTACATGATCCCCTGTGAGAAGGTGTCCACCCTGCCCGCGATCACACTGAAGCTGGGAGGCAAAGGCT
CTSD.2	80	ACAAGCTGTCCC
		GCAAGTTCCAACCTGGAAAGGCCATCGGCTTTGTCAAGGATGTAGCCAACATCACAATCTATGACGAGG
CTSH.2	77	AAGCGATG
		GGGAGGCCTTATCTCACTGAGTGAGCAGAATCTGGTAGACTGCTCTGGGCCTCAAGGCAATGAAGGCTGC
CTSL.2	74	AATGG
CTSL2.1	67	TGTCTCACTGAGCGAGCAGAATCTGGTGGACTGTTTCGCGTCCTCAAGGCAATCAGGGCTGCAATGGT
		TGACAACGGCTTTCCAGTACATCATTGATAACAAGGGCATCGACTCAGACGCTTCCTATCCCTACAAAGC
CTSS.1	76	CATGGA
		GAGGCCGTTACTGTGGCACCGACATGCCCCATCCTATCACATCCTTCAGCAGCGCCCTGACGCTGAGAT
CUBN.1	71	TC
		ATGCCCTGGTAATGTCTGCATTCAACAATGACGCTGGCTTTGTGGCTGCTCTTGATAAGGCTTGTGGTGC
CUL1.1	71	C
		AAGCATCTTCCTGTTCTTGGACCGCACCTATGTGCTGCAGAACTCCACGCTGCCCTCCATCTGGGATAT
CUL4A.1	75	GGGATT

Table B

Gene	Target Sequence Length	Amplicon Sequence
CX3CL1.1	66	GACCCCTTGCCGTCTACCTGAGGGGCCTCTTATGGGCTGGGTTCTACCCAGGTGCTAGGAACACTCC
CX3CR1.1	68	TTCCCAGTTGTGACATGAGGAAGGATCTGAGGCTGGCCCTCAGTGTGACTGAGACGGTTGCATTTAGC
CXCL10.1	68	GGAGCAAATCGATGCAGTGCTTCCAAGGATGGACCACACAGAGGCTGCCTCTCCCATCACTTCCCTA
CXCL12.1	67	GAGCTACAGATGCCCATGCCGATTCTTCGAAAGCCATGTTGCCAGAGCCAACGTCAAGCATCTCAAA
CXCL14.1	74	TGCGCCCTTTCTCTGTACATATACCCTTAAGAACGCCCCCTCCACACACTGCCCCCAGTATATGCCCGC
CXCL9.1	70	ATTG ACCAGACCATTGTCTCAGAGCAGGTGCTGGCTCTTTCTGGCTACTCCATGTTGGCTAGCCTCTGGTAA
CXCR4.3	72	C TGACCGCTTCTACCCCAATGACTTGTGGGTGGTTGTGTTCCAGTTTCAGCACATCATGGTTGGCCTTATC
CXCR6.1	67	CT CAGAGCCTGACGGATGTGTTCTGGTGAACCTACCCCTGGCTGACCTGGTGTGTTGTCTGCACTCTGC
CYP2C8.2	73	CCGTGTTCAAGAGGAAGCTCACTGCCTTGTGGAGGAGTTGAGAAAAACCAAGGCTTCACCCTGTGATCC
CYP2C8v2.1	70	CACT GCTGTAGTGACGAGATCCAGAGATACAGTGACCTTGTCACCCACCGGTGTGCCCCATGCAGTGACCACT
CYP3A4.2	79	G AGAACAAGGACAACATAGATCCTTACATATACACACCCTTTGGAAGTGGACCCAGAACTGCATTGGCAT
CYR61.1	76	GAGGTTTGC TGCTCATTCTTGAGGAGCATTAAAGGTATTTGAAACTGCCAAGGGTGCTGGTGCGGATGGACACTAATG
DAG1.1	67	CAGCCAC GTGACTGGGCTCATGCCTCCAAGTCAGAGTTTCCCTGGTGCCCCAGAGACAGGAGCACAAGTGGGAT
DAPK1.3	77	CGCTGACATCATGAATGTTCTCGACCGGCTGGAGGCGAGTTTGGATATGACAAAGACACATCGTTGCT
DCBLD2.1	69	GAAAGAGA TCACCAGGGCAGGAAGTTTATCATGCCTATGCTGAACCACTCCCAATTACGGGGCCTGAGTATGCAACC
DCC.3	75	AAATGTCTCCTCGACTGCTCCGCGGAGTCCGACCGAGGAGTTCCAGTGATCAAGTGGAAGAAAGATGG
DCN.1	67	CATTCA GAAGGCCACTATCATCCTCCTTCTGCTTGACACAAGTTTCTGGGCTGGACCGTTTCAACAGAGAGGC
DCXR.1	66	CCATAGCGTCTACTGCTCCACCAAGGGTGCCCTGGACATGCTGACCAAGGTGATGGCCCTAGAGCT
DDC.1	67	CAGAGCCCAGACACCATGAACGCAAGTGAATTCGAAGGAGAGGGAAGGAGATGGTGGATTACGTGG
DEFB1.1	68	GATGGCCTCAGGTGGTAACCTTCTCACAGGCCTTGCCACAGATCTGATCATTACAATTGCGTCAGCA

Table B

Gene	Target Sequence Length	Amplicon Sequence
DET1.1	70	CTTGTGGAGATCACCCAATCAGGTTCTATGCCCCGGGACTCGGGCCTGCTCAAGTTTGAGATCCAGGCGG G
DHPS.3	78	GGGAGAACGGGATCAATAGGATCGGAAACCTGCTGGTGCCCAATGAGAATTACTGCAAGTTTGAGGACT GGCTGATGC
DIABLO.1	73	CACAATGGCGGCTCTGAAGAGTTGGCTGTGCGCGAGCGTAACTTCATTCTTCAGGTACAGACAGTGTTT GTGT
DIAPH1.1	62	CAAGCAGTCAAGGAGAACCAGAAGCGGCGGGAGACAGAAGAAAAGATGAGGCGAGCAAACT
DICER1.2	68	TCCAATTCCAGCATCACTGTGGAGAAAAGCTGTTTGTCTCCCCAGCATACTTTATCGCCTTCACTGCC
DKFZP564O0823.1	66	CAGCTACACTGTGCGAGTCCGCTGCTGAGCCTCCACACTCATCTCCCCTCAAGCTCCAGCCTCAT
DLC1.1	68	GATTTCAGACGAGGATGAGCCTTGTGCCATCAGTGGCAAATGGACTTTCCAAAGGGACAGCAAGAGGTG
DLL4.1	67	CACGGAGGTATAAGGCAGGAGCCTACCTGGACATCCCTGCTCAGCCCCGCGGCTGGACCTTCCTTCT GGACTCCAGATGCCAGGAGCCCTGCTGCCCACATGCAAGGACCAGCATCTCCTGAGAGGACGCCTGGG
DPEP1.1	72	CTTA
DPYS.1	70	AAAGAATGGCACCATGCAGCCCACCATGTGATGGGTCCACCTTTGCGACCAGACCCCTCAACACCCGAC T
DR4.2	83	TGCACAGAGGGTGTGGGTTACACCAATGCTTCCAACAATTTGTTTGCTTGCCTCCCATGTACAGCTTGTA AATCAGATGAAGA
DR5.2	84	CTCTGAGACAGTGCTTCGATGACTTTGCAGACTTGGTGCCCTTTGACTCCTGGGAGCCGCTCATGAGGA AGTTGGGCCTCATGG
DUSP1.1	76	AGACATCAGCTCCTGGTTCAACGAGGCCATTGACTTCATAGACTCCATCAAGAATGCTGGAGGAAGGGT GTTTGTC
DUSP9.1	77	CGTCCTAATCAACGTGCCTATGGCGGGACCACGCTCGGAGCCTGCCTCTTCTGCGACTGTTACTTTTTT TTTGCGGG
E2F1.3	75	ACTCCCTCTACCCTTGAGCAAGGGCAGGGGTCCCTGAGCTGTTCTTCTGCCCCATACTGAAGGAACTGA GGCCTG
EBAG9.1	66	CGCTCCTGTTTTTCTCATCTGTGCAGTGGGTTTTGATTCCCACCATGGCCATCACCCAGTTTCGGT
ECRG4.1	66	GCTCCTGCTCCTGTGCTGGGGCCCAGGTGGCATAAGTGGAATAAACTCAAGCTGATGCTTCAAAA ACGAGTCCATTGCCTTCTTTTATAACCGAAGTGGAAGCATCTTGCCACAGAATGGAACACAGTCAGCAA
EDG2.1	72	GC
EDN1 endothelin.1	73	TGCCACCTGGACATCATTTGGGTCAACACTCCCGAGCACGTTGTTCCGTATGGACTTGGAAGCCCTAGG TCCA
EDN2.1	79	CGACAAGGAGTGCGTCTACTTCTGCCACTTGGACATCATCTGGGTGAACACTCCTGAACAGACAGCTCC TTACGGCCTG

Table B

Gene	Target Sequence Length	Amplicon Sequence
EDNRA.2	76	TTTCCTCAAATTTGCCTCAAGATGGAAACCCTTTGCCTCAGGGCATCCTTTTGGCTGGCACTGGTTGGAT GTGTAA
EDNRB.1	72	ACTGTGAACTGCCTGGTGCAGTGTCCACATGACAAAGGGGCAGGTAGCACCTCTCTCACCCATGCTGT GGT
EEF1A1.1	67	CGAGTGGAGACTGGTGTCTCAAACCCGGTATGGTGGTCACCTTTGCTCCAGTCAACGTTACAACGG
EFNB1.2	66	GGAGCCCGTATCCTGGAGCTCCCTCAACCCCAAGTTCCTGAGTGGGAAGGGCTTGGTGATCTATCC TGACATTATCATCCCGCTAAGGACTGCGGACAGCGTCTTCTGCCCTCACTACGAGAAGGTCAGCGGGGA
EFNB2.1	73	CTAC CTTTGCCTTGCTCTGTGCACAGTGAAGTCAGCCAGAGCAGGGCTGTAAACTCTGTGAAATTTGTCATAAG
EGF.3	84	GGTGTCAGGTATTT
EGFR.2	62	TGTCGATGGACTTCCAGAACCACCTGGGCAGCTGCCAAAAGTGTGATCCAAGCTGTCCCAAT
EGLN3.1	68	GCTGGTCCTCTACTGCGGGAGCCGGCTGGGCAAATACTACGTCAAGGAGAGGTCTAAGGCAATGGTGG GTCCCCGCTGCAGATCTCTGACCCGTTGGATCCTTTCCTCACTCGCCACCATGGACAACCTACCCTAA
EGR1.1	76	GCTGGAG
EIF2C1.1	67	CCCTCACGGACTCTCAGCGCGTTTCGCTTCACCAAGGAGATCAAGGGCCTGAAGGTGGAAGTCACCCA
EIF4EBP1.1	66	GGCGGTGAAGAGTCACAGTTTGAGATGGACATTTAAAGCACCAGCCATCGTGTGGAGCACTACCAA
ELTD1.1	66	AGGTCTTGTCGAAGAGGAGCCCTCGCTCTTCTGTTCTTCTCGGCACCACCTGGATCTTTGGGGTT AGGCACTGAGGGTGGAAAAAATGCAAGCACTTCAGCAACCAGCCGGTCTTATTCAGTATTATTTTGCCG
EMCN.1	73	GTG GCTAGTACTTTGATGCTCCCTTGATGGGGTCCAGAGAGCCTCCCTGCAGCCACCAGACTTGGCCTCCAG
EMP1.1	75	CTGTTG
ENO2.1	67	TCCTTGGCTTACCTGACCTCTTGCTGTCTCTGCTCGCCCTCCTTTCTGTGCCCTACTCATTGGGGTT
ENPEP.1	67	CACCTACACGGAGAACGGACAAGTCAAGAGCATAGTGGCCACCGATCATGAACCAACAGATGCCAGG
ENPP2.1	67	CTCCTGCGCACTAATACCTTCAGGCCAACCATGCCAGAGGAAGTTACCAGACCCAATTATCCAGGGA AAGCCTTGGAGGGTTTCATTGCCGTGGTGACCCAAGATGGCGACATGATCTTCTGTGAGAAAACATCA
EPAS1.1	72	GCA
EPB41L3.1	66	TCAGTGCCATACGCTCTCACTCTCTCCTTCCCTCTGGCTCTGTGCCTCTGCTACCTGGAGCCCAAG CGCCTGTTCAACCAAGATTGACACCATTGCGCCCGATGAGATCACCGTCAGCAGCGACTTCGAGGCACGC
EPHA2.1	72	CAC
EPHB1.3	67	CCTTGGGAGGGAAGATCCCTGTGAGATGGACAGCTCCAGAGGCCATCGCCTACCGCAAGTTCACTTC

Table B

Gene	Target Sequence Length	Amplicon Sequence
EPHB2.1	66	CAACCAGGCAGCTCCATCGGCAGTGTCCATCATGCATCAGGTGAGCCGCACCGTGGACAGCATTAC
EPHB4.1	77	TGAACGGGGTATCCTCCTTAGCCACGGGGCCCCGTCCCATTGAGCCTGTCAATGTCACCACTGACCGAG
EPO.1	84	AGGTACCT CAGTGCCAGCAATGACATCTCAGGGGCCAGAGGAACTGTCCAGAGAGCAACTCTGAGATCTAAGGATGT
ErbB3.1	81	CACAGGGCCAACTTG CGGTTATGTCATGCCAGATACACACCTCAAAGGTA CTCCCTCCTCCCGGGAAGGCACCCCTTTCTTCAGT
ERBB4.3	86	GGGTCTCAGTTC TGGCTCTTAATCAGTTTCGTTACCTGCCTCTGGAGAATTTACGCATTATTCGTGGGACAAAACCTTTATGAG
ERCC1.2	67	GATCGATATGCCTTG GTCCAGGTGGATGTGAAAGATCCCCAGCAGGCCCTCAAGGAGCTGGCTAAGATGTGTATCCTGGCCG
ERCC4.1	67	CTGCTGGAGTACGAGCGACAGCTGGTGCTGGAAC TGTCTCGACACTGACGGGCTAGTAGTGTGCGCCC
EREG.1	91	ATAACAAAGTGTAGCTCTGACATGAATGGCTATTGTTTGCATGGACAGTGCATCTATCTGGTGGACATGA
ERG.1	70	GTCAAAACTACTGCAGGTGTG CCAACACTAGGCTCCCCACCAGCCATATGCCTTCTCATCTGGGCACTTACTACTAAAGACCTGGCGGAG
ERK1.3	67	G ACGGATCACAGTGGAGGAAGCGCTGGCTCACCCCTACCTGGAGCAGTACTATGACCCGACGGATGAG
ERK2.3	68	AGTTCTTGACCCCTGGTCCTGTCTCCAGCCCGTCTTGGCTTATCCACTTTGACTCCTTTGAGCCGTTT
ESPL1.3	70	ACCCCCAGACCGGATCAGGCAAGCTGGCCCTCATGTCCCCTTCACGGTGTTTGAGGAAGTCTGCCCTAC
ESRRG.3	67	A CCAGCACCATTTGTTGAAGATCCCCAGACCAAGTGTGAATACATGCTCAACTCGATGCCCAAGAGACT
F2.1	77	GCTGCATGTCTGGAAGGTA ACTGTGCTGAGGGTCTGGGTACGA ACTACCGAGGGCATGTGAACATCACC
F3.1	73	CGGTCAGG GTGAAGGATGTGAAGCAGACGTACTTGGCACGGGTCTTCTCCTACCCGGCAGGGAATGTGGAGAGCAC
FABP1.1	66	CGGT GGGTCCAAAGTGATCCAAAACGAATTCACGGTGGGGGAGGAATGTGAGCTGGAGACAATGACAGGG
FABP7.1	72	GGAGACAAAGTGGTCATCAGGACTCTCAGCACATTCAAGAACACGGAGATTAGTTTCCAGCTGGGAGAA
FAP.1	66	GAG CTGACCAGAACACGGCTTATCCGGCCTGTCCACGAACCACTTATACACCCACATGACCCACTTCC
fas.1	91	GGATTGCTCAACAACCATGCTGGGCATCTGGACCCTCCTACCTCTGGTTCTTACGTCTGTTGCTAGATTA
		TCGTCCAAAAGTGTTAATGCC

Table B

Gene	Target Sequence Length	Amplicon Sequence
fasl.2	80	GCACTTTGGGATTCTTTCCATTATGATTCTTTGTTACAGGCACCGAGAATGTTGTATTCAGTGAGGGTCTT CTTACATGC
FBXW7.1	73	CCCCAGTTTCAACGAGACTTCATTTCAATTGCTCCCTAAAGAGTTGGCACTCTATGTGCTTTCATTCTGGA AC
FCER1G.2	73	TGCCATCCTGTTTCTGTATGGAATTGTCCTCACCTCCTCTACTGTGCGACTGAAGATCCAAGTGCGAAAG GCA
FCGR3A.1	67	GTCTCCAGTGGAAGGGAAAAGCCCATGATCTTCAAGCAGGGAAGCCCCAGTGAGTAGCTGCATTCCT GGATGATTACCTTGACCTCTTTGGGGACCCAGTGTGACCGGCAAATTGGCACTGACATCCAGGACAA
FDPS.1	77	CAAATGCA
FEN1.1	66	GTGGAGAAGGGTACGCCAGGGTGCCTGAGAGACTCTGTTCTCCCTGGAGGGACTGGTTGCCATGAG
FGF1.1	66	GACACCGACGGGCTTTTATACGGCTCACAGACACCAAATGAGGAATGTTTGTTCCTGGAAGGCTG AGATGCAGGAGAGAGGAAGCCTTGCAAACCTGCAGACTGCTTTTTGCCCAATATAGATTGGGTAAAGGCT
FGF2.2	76	GCAAAAC
FGF9.1	67	CACAGCTGCCATACTTCGACTTATCAGGATTCTGGCTGGTGGCCTGCGCGAGGGTGCAGTCTTACTT CACGGGACATTCACCACATCGACTACTATAAAAAGACAACCAACGGCCGACTGCCTGTGAAGTGGATGG
FGFR1.3	74	CACCC GAGGGACTGTTGGCATGCAGTGCCCTCCCAGAGACCAACGTTCAAGCAGTTGGTAGAAGACTTGGATCG
FGFR2 isoform 1.1	80	AATTCTCACTC
FH.1	67	ATGGTTGCAGCCCAAGTCATGGGGAACCATGTTGCTGTCACTGTCGGAGGCAGCAATGGACATTTTG
FHIT.1	67	CCAGTGGAGCGCTTCCATGACCTGCGTCCTGATGAAGTGGCCGATTTGTTTCAGACGACCCAGAGAG
FHL1.1	66	ATCCAGCCTTTGCCGAATACATCCTATCTGCCACACATCCAGCGTGAGGTCCCTCCAGCTACAAGG GGTTCCAGCTTTCTGTAGCTGTAAGCATTGGTGGCCACACCACCTCCTTACAAAGCAACTAGAACCTGCG
FIGF.1	72	GC
FILIP1.1	66	ACACCGGTCACAACGTCATCTGCTCGAGGAACCCAGTCAGTGTGAGGACAAGACGGGTCATCCCAG CTGCCCTGACTGAATGTGTTCTGTCACTCAGCTTGCTTCCGACACCTCTGTTTCCTCTTCCCCTTTCTC
FKBP1A.1	76	CTCGTA
FLJ22655.1	82	CTCCTTCACACAGAACCTTTTCAATTTATTGTACAACATCACACTCACCTAACCTACTGGCGGACAGCGATC CCAGTTTGCCT
FLT1.1	75	GGCTCCTGAATCTATCTTTGACAAAATCTACAGCACCAAGAGCGACGTGTGGTCTTACGGAGTATTGCTG TGGGA
FLT3LG.1	71	TGGGTCCAAGATGCAAGGCTTGCTGGAGCGCGTGAACACGGAGATACACTTTGTACCAAATGTGCCTT TC

Table B

Gene	Target Sequence Length	Amplicon Sequence
FLT4.1	69	ACCAAGAAGCTGAGGACCTGTGGCTGAGCCCGCTGACCATGGAAGATCTTGTCTGCTACAGCTTCCAGG
FN1.1	69	GGAAGTGACAGACGTGAAGGTCACCATCATGTGGACACCGCCTGAGAGTGCAGTGACCGGCTACCGTG T
FOLR1.1	67	GAACGCCAAGCACCACAAGGAAAAGCCAGGCCCGAGGACAAGTTGCATGAGCAGTGTGACCCCTGG
FOS.1	67	CGAGCCCTTTGATGACTTCCTGTTCCAGCATCATCCAGGCCAGTGGCTCTGAGACAGCCCGCTCC
FRAP1.1	66	AGCGCTAGAGACTGTGGACCGCCTGACGGAGTCCCTGGATTTCACTGACTATGCCTCCCGGATCAT TTGGTACCTGTGGGTTAGCATCAAGTTCTCCCCAGGGTAGAATTCAATCAGAGCTCCAGTTTGCATTTGG
FRP1.3	75	ATGTG GTAAGTCGGATGAGCCTGTCTGTGCCAGTGACAATGCCACTTATGCCAGCGAGTGTGCCATGAAGGAAG
FST.1	72	CTG TGGATCCTCACCTGGTCCGGTGCTGTGCTGCGCTTCCACCTTCTTCACTGTCACCACGTA CTTGGTAGAC
FZD2.2	78	ATGCAGCGC
G-Catenin.1	68	TCAGCAGCAAGGGCATCATGGAGGAGGATGAGGCCTGCGGGCGCCAGTACACGCTCAAGAAAACCACC
GADD45B.1	70	ACCCTCGACAAGACCACACTTTGGGACTTGGGAGCTGGGGCTGAAGTTGCTCTGTACCCATGA ACTCCC A
GAS2.1	68	AACATGTCATGGTCCGTGTGGGAGGAGGCTGGGAACTTTTGCAGGGTATTTGTTGAAACACGACCCC
GATA3.3	75	CAAAGGAGCTCACTGTGGTGTCTGTGTTCCAACCACTGAATCTGGACCCCATCTGTGAATAAGCCATTCT GACTC
GATM.1	67	GATCTCGGCTTGGACGAACCTTGACAGGATGGGTGCAGCGAACTTTCCAGAGCACCCAGGCAGCTAC
GBL.1	66	GCTGTCAATAGCACCGGAACTGCTATGTCTGGAATCTGACGGGGGGCATTGGTGACGAGGTGACC GCATGGGAACCATCAACCAGCAGGCCATGGACCAACTTCACTATGTGACAGAGCTGACAGATCGAATCA
GBP2.2	83	AGGCAAACCTCCTCA CTGTTGCAGGAAGGCATTGATCATCTCCTGGCCCAGCATGTTGCTCATCTCTTTATTAGAGACCCACTGA
GCLC.3	71	C TGTAGAATCAAACCTTTCATCATCAACTAGAAAGTGCAGTTGACATGGCCTGTTCACTCCTGGAGTTGCA
GCLM.2	85	CAGCTGGATTCTGTG
GFRA1.1	69	TCCGGGTAAAGAACAAGCCCCTGGGGCCAGCAGGGTCTGAGAATGAAATTCCCACTCATGTTTTGCCAC

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Gene	Target Sequence Length	Amplicon Sequence
GJA1.1	68	GTTCACTGGGGGTGTATGGGGTAGATGGGTGGAGAGGGAGGGGATAAGAGAGGTGCATGTTGGTATTT
GLYAT.1	68	TACCATTGCAAGGTGCCCAGATGCTGCAGATGCTGGAGAAATCCTTGAGGAAGAGCCTCCCAGCATCC
GMNN.1	67	GTTCGCTACGAGGATTGAGCGTCTCCACCCAGTAAGTGGGCAAGAGGCGGCAGGAAGTGGGTACGCA
GNAS.1	72	GAACGTGCCTGACTTTGACTTCCCTCCCGAATTCTATGAGCATGCCAAGGCTCTGTGGGAGGATGAAGG AGT
GPC3.1	68	TGATGCGCCTGGAAACAGTCAGCAGGCAACTCCGAAGGACAACGAGATAAGCACCTTTCACAACCTCG
GPX1.2	67	GCTTATGACCGACCCCAAGCTCATCACCTGGTCTCCGGTGTGTGCGCAACGATGTTGCCTGGAACCTT
GPX2.2	75	CACACAGATCTCCTACTCCATCCAGTCTGAGGAGCCTTAGGATGCAGCATGCCTTCAGGAGACACTGC TGGACC
GPX3.1	69	GCTCTAGGTCCAATTGTTCTGCTCTAACTGATACCTCAACCTTGGGGCCAGCATCTCCCACTGCCTCCA
GRB14.1	76	TCCCACTGAAGCCCTTTCAGTTGCGGTTGAAGAAGGACTCGCTTGGAGGAAAAAGGATGTTTACGCCT GGGCACT
GRB7.2	67	CCATCTGCATCCATCTTGTTTGGGCTCCCCACCCTTGAGAAGTGCCTCAGATAATACCCTGGTGGCC
GRO1.2	73	CGAAAAGATGCTGAACAGTGACAAATCCAAGTACCAGAAGGGAGGAGGAAGCTCACTGGTGGCTGTTC CTGA
GSTM1.1	86	AAGCTATGAGGAAAAGAAGTACACGATGGGGGACGCTCCTGATTATGACAGAAGCCAGTGGCTGAATGA AAAATTCAAGCTGGGCC
GSTM3.2	76	CAATGCCATCTTGCGCTACATCGCTCGCAAGCACAACATGTGTGGTGAGACTGAAGAAGAAAAGATTCCG AGTGGAC
GSTp.3	76	GAGACCCTGCTGTCCCAGAACCAGGGAGGCAAGACCTTCATTGTGGGAGACCAGATCTCCTTCGCTGAC TACAACC
GSTT1.3	66	CACCATCCCCACCCTGTCTTCCACAGCCGCCTGAAAGCCACAATGAGAATGATGCACACTGAGGCC
GZMA.1	79	GAAAGAGTTTCCCTATCCATGCTATGACCCAGCCACACGCGAAGGTGACCTTAACTTTTACAGCTGACG GAAAAAGCA
HADH.1	66	CCACCAGACAAGACCGATTGCGTGGCCTCCATTTCTTCAACCCAGTGCCTGTCATGAACTTGTGG
HAVCR1.1	76	CCACCAAGGTCACGACTACTCCAATTGTCAAACTGTTCCAACCGTCACGACTGTTTGAACGAGCACCA CTGTTC
HDAC1.1	74	CAAGTACCACAGCGATGACTACATTAATTCTTGCGCTCCATCCGTCCAGATAACATGTCTGGAGTACAGC AAGC

Table B

Gene	Target Sequence Length	Amplicon Sequence
Hepsin.1	84	AGGCTGCTGGAGGTCATCTCCGTGTGTGATTGCCCCAGAGGCCGTTTCTTGGCCGCCATCTGCCAAGA CTGTGGCCGCAGGAAG
HER2.3	70	CGGTGTGAGAAGTGCAGCAAGCCCTGTGCCCGAGTGTGCTATGGTCTGGGCATGGAGCACTTGCGAGA GG
HGD.1	76	CTCAGGTCTGCCCCTACAATCTCTATGCTGAGCAGCTCTCAGGATCGGCTTTCAC TTGTCCACGGAGCA CCAATAA
HGF.4	65	CCGAAATCCAGATGATGATGCTCATGGACCCTGGTGCTACACGGGAAATCCACTCATTCTTGGG CAGGACACAAGTGCCAGATTGCGGGCTGGGGCCACTTGATGAGAACGTGAGCGGCTACTCCAGCTCC
HGFAC.1	72	CTGC TGAACATAAAGTCTGCAACATGGAAGGTATTGCACTGCACAGGCCACATTCACGTATATGATACCAACAG
HIF1A.3	82	TAACCAACCTCA
HIF1AN.1	66	TGTTGGCCAGGTCTCACTGCAGCCTGCCCCAGGCTAACTGGCTAGAGCCTCCAGGCCCTATGATGC
HIST1H1D.1	67	AAAAAGGCCGAAGAAGGCAGGCGCAACTGCTGGGAAACGCAAAGCATCCGGACCCCCAGTATCTGAGC CTTGTGAGGGACTGAGATGCAGGATTTCTTCACGCCTCCCCTTTGTGACTTCAAGAGCCTCTGGCATCT
HLA-B.1	78	CTTTCTGCA CGCCCTGAAGACAGAATGTTCCATATCAGAGCTGTGATCTTGAGAGCCCTCTCCTTGGCTTTCCTGCTGA
HLA-DPA1.1	78	GTCTCCGA TCCATGATGGTTCTGCAGGTTTCTGCGGCCCCCGGACAGTGGCTCTGACGGCGTTACTGATGGTGCT
HLA-DPB1.1	73	GCTCA
HLA-DQB1.1	67	GGTCTGCTCGGTGACAGATTTCTATCCAGGCCAGATCAAAGTCCGGTGGTTTTCGGAATGATCAGGAG CATCTTTCTCCTGTGGTCAACATCACATGGCTGAGCAATGGGCAGTCAGTCACAGAAGGTGTTTCTGA
HLADQA1.2	76	GACCAGC TGGCCTGTCCATTGGTGATGTTGCGAAGAACTGGGAGAGATGTGGAATAACACTGCTGCAGATGACAA
HMGB1.1	71	GC AGCAGGAGCGACCAACTGATCGCACACATGCTTTGTTTGGATATGGAGTGAACACAATTATGTACCAAAT
HNRPAB.1	84	TTAACTTGGCAAAC CAGGCAGATGGACACCAACAATGACGGCAAACGTGCTTGGAGAATTTCATCAGAGGTGCCAAGAGCGA
HPCAL1.1	70	C AGCTGAAGACGGCCAAGATCAAGGTGAAGGAGAACATTGATGCCCTGGAGGAGCTGAAAATCCTGGTG
HPD.1	78	GACTACGACG
HSD11B2.1	69	CCAACCTGCCTCAAGAGCTGCTGCAGGCCTACGGCAAGGACTACATCGAGCACTTGATGGGCAGTTCC
HSP90AB1.1	66	GCATTGTGACCAGCACCTACGGCTGGACAGCCAATATGGAGCGGATCATGAAAGCCCAGGCACTTC

Table B

Gene	Target Sequence Length	Amplicon Sequence
HSPA1A.1	70	CTGCTGCGACAGTCCACTACCTTTTTCGAGAGTGA CTCCCGTTGTCCCAAGGCTTCCAGAGCGAACCTG
HSPA8.1	73	CCTCCCTCTGGTGGTGTCTCCTCAGGGCCACCATTGAAGAGGTTGATTAAGCCAACCAAGTGTAGATGTAGC
HSPB1.1	84	CCGACTGGAGGAGCATAAAAGCGCAGCCGAGCCCAGCGCCCCGCACTTTTCTGAGCAGACGTCCAGAGCAGAGTCAGCCAGCAT
HSPG2.1	66	GAGTACGTGTGCCGAGTGTTGGGCAGCTCCGTGCCTCTAGAGGCCTCTGTCCTGGTCACCATTGAG
HTATIP.1	66	TCTGAATTGTTTGGGCACTGATGAGGACTCCAGGACAGCTCTGATGGAATACCGTCAGCACCACGC
		TGGCTGTGGAGTTCAAATGTCTGATGCTACCCTGGCTGGCAGGCACCGTGGTGTGAGCGGAAGAGCATGTGGTGATTGG
HYAL1.1	78	TGGTGATTGG
HYAL2.1	67	CAACCATGCACTCCCAGTCTACGTCTTCACACGACCCACCTACAGCCGCAGGCTCACGGGGCTTAGT
HYAL3.1	67	TATGTCCGCCTCACACACCGGAGATCTGGGAGGTTCTGTCCCAGGATGACCTTGTGCAGTCCATTGTATGTCCGCCTCACACACCGGAGATCTGGGAGGTTCTGTCCCAGGATGACCTTGTGCAGTCCATTG
ICAM1.1	68	GCAGACAGTGACCATCTACAGCTTTCCGGCGCCCCAACGTGATTCTGACGAAGCCAGAGGTCTCAGAAG
ICAM2.1	62	GGTCATCCTGACACTGCAACCCACTTTGGTGGCTGTGGGCAAGTCCTTCACCATTGAGTGCA
ICAM3.1	67	GCCTTCAATCTCAGCAACGTGACTGGCAACAGTCGGATCCTCTGCTCAGTGTACTGCAATGGCTCTC
		AGAACCGCAAGGTGAGCAAGGTGGAGATTCTCCAGCACGTCATCGACTACATCAGGGACCTTCAGTTGGAACCGCAAGGTGAGCAAGGTGGAGATTCTCCAGCACGTCATCGACTACATCAGGGACCTTCAGTTGG
ID1.1	70	A
		AACGACTGCTACTCCAAGCTCAAGGAGCTGGTGCCCAGCATCCCCCAGAACAAGAAGGTGAGCAAGATGAACGACTGCTACTCCAAGCTCAAGGAGCTGGTGCCCAGCATCCCCCAGAACAAGAAGGTGAGCAAGATG
ID2.4	76	GAAATCC
		CTTCACCAAATCCCTTCCTGGAGACTAAACCTGGTGCTCAGGAGCGAAGGACTGTGAACTTGTGGCCTGAACGACTGCTACTCCAAGCTCAAGGAGCTGGTGCCCAGCATCCCCCAGAACAAGAAGGTGAGCAAGATG
ID3.1	80	AAGAGCCAGAG
		CTCTCCGGATTGACCAAGTTCATCCTGGGCTCCATTGGGTCTGCCATTGCGGCTGTCATTGCGAGGTTC
IFI27.1	71	TA
		TCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTATTGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCT
IGF1.2	76	CTGTCCG
		GCATGGTAGCCGAAGATTTACAGTCAAAATCGGAGATTTTGGTATGACGCGAGATATCTATGAGACAGACTATTACCGGAAA
IGF1R.3	83	CTATTACCGGAAA
		CCGTGCTTCCGGACAACCTCCCCAGATACCCCGTGGGCAAGTTCTTCCAATATGACACCTGGAAGCAGT
IGF2.2	72	CCA
		GTGGACAGCACCATGAACATGTTGGGCGGGGGAGGCAGTGCTGGCCGGAAGCCCCTCAAGTCGGGTA
IGFBP2.1	73	TGAAGG
IGFBP3.1	66	ACATCCCAACGCATGCTCCTGGAGCTCACAGCCTTCTGTGGTGTCAATTTCTGAAACAAGGGCGTGG

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Gene	Target Sequence Length	Amplicon Sequence
IGFBP5.1	69	TGGACAAGTACGGGATGAAGCTGCCAGGCATGGAGTACGTTGACGGGGACTTTCAGTGCCACACCTTC G
IGFBP6.1	77	TGAACCGCAGAGACCAACAGAGGAATCCAGGCACCTCTACCACGCCCTCCCAGCCCAATTCTGCGGGT GTCCAAGAC
IL-7.1	71	GCGGTGATTTCGAAATTCGCGAATTCCTCTGGTCCTCATCCAGGTGCGCGGGAAGCAGGTGCCAGGA GAG
IL-8.1	70	AAGGAACCATCTCACTGTGTGTAAACATGACTTCCAAGCTGGCCGTGGCTCTCTTGGCAGCCTTCCTGAT GGCGCTGTCTCGATTTCTTCCCTGTGAAAACAAGAGCAAGGCCGTGGAGCAGGTGAAGAATGCCTTTA
IL10.3	79	ATAAGCTCCA
IL11.2	66	TGGAAGGTTCCACAAGTCACCCTGTGATCAACAGTACCCGTATGGGACAAAGCTGCAAGGTCAAGA GGCTGGGTACCAATGCTGCAGGTCAACAGCTATGCTGGTAGGCTCCTGCCAGTGTGGAACCACTGACTA
IL15.1	79	CTGGCTCTCA
IL1B.1	67	AGCTGAGGAAGATGCTGGTTCCTGCCACAGACCTTCCAGGAGAATGACCTGAGCACCTTCTTTCC CCTGAACCTTCCAAAGATGGCTGAAAAAGATGGATGCTTCCAATCTGGATTCAATGAGGAGACTTGCCTG
IL6.3	72	GT
IL6ST.3	74	GGCCTAATGTTCCAGATCCTTCAAAGAGTCATATTGCCAGTGGTCACCTCACACTCCTCCAAGGCACAA TTTT
ILT-2.2	63	AGCCATCACTCTCAGTGCGAGCCAGGTCCTATCGTGGCCCCTGAGGAGACCCTGACTCTGCAGT GTGGACTCGTCCAAGATCAAGCGGCACGTGCTAGAGTACAATGAGGAGCGCGATGACTTCGATCTGGAA
IMP3.1	72	GCC
INDO.1	66	CGCCTTGCACGTCTAGTTCTGGGATGCATCACCATGGCATATGTGTGGGGCAAAGGTCATGGAGAT GTGCCCCGAGCCATATAGCAGGCACGTCCGGGTCCTCACTGTCCTTCCACTCAACAGTCATCAACCACTA
INHBA.1	72	CCG
INHBB.1	72	AGCCTCCAGGATACCAGCAAATGGATGCGGTGACAAATGGCAGCTTAGCTACAAATGCCTGTCAGTCGG AGA
INSR.1	67	CAGTCTCCGAGAGCGGATTGAGTTCCTCAATGAGGCCTCGGTCATGAAGGGCTTCACCTGCCATCAC
IQGAP2.1	66	AGAGACACCAGCAACTGCGCAACAGGAGGTAGACCATGCCACGGACATGGTGAGCCGTGCAATGAT GTGTCAGACTGAAGCCCCATCCAGCCCGTTCCGCAGGGACTAGAGGCTTTTCGGCTTTTTGGGACAGCAA
ISG20.1	70	C
ITGA3.2	77	CCATGATCCTCACTCTGCTGGTGGACTATACACTCCAGACCTCGCTTAGCATGGTAAATCACCGGCTACA AAGCTTC
ITGA4.2	66	CAACGCTTCAGTGATCAATCCCGGGGCGATTACAGATGCAGGATCGGAAAGAATCCCGGCCAGAC

Table B

Gene	Target Sequence Length	Amplicon Sequence
ITGA5.1	75	AGGCCAGCCCTACATTATCAGAGCAAGAGCCGGATAGAGGACAAGGCTCAGATCTTGCTGGACTGTGGA GAAGAC
ITGA6.2	69	CAGTGACAAACAGCCCTTCCAACCCAAGGAATCCCACAAAAGATGGCGATGACGCCCATGAGGCTAAAC GATATGATTGGTCGCTGCTTTGTGCTCAGCCAGGACCTGGCCATCCGGGATGAGTTGGATGGTGGGGA
ITGA7.1	79	ATGGAAGTTCT ACTCGGACTGCACAAGCTATTTTTGATGACAGCTATTTGGGTATTCTGTGGCTGTCTGGAGATTTCATG
ITGAV.1	79	GTGATGGCA TCAGAATTGGATTTGGCTCATTTGTGGAAAAGACTGTGATGCCTTACATTAGCACAACACCAGCTAAGCT
ITGB1.1	74	CAGG ACCGGGGAGCCCTACATGACGAAAATACCTGCAACCGTTACTGCCGTGACGAGATTGAGTCAGTGAAAG
ITGB3.1	78	AGCTTAAGG
ITGB4.2	66	CAAGGTGCCCTCAGTGGAGCTCACCAACCTGTACCCGTATTGCGACTATGAGATGAAGGTGTGCGC TCGTGAAAGATGACCAGGAGGCTGTGCTATGTTTCTACAAAACCGCCAAGGACTGCGTCATGATGTTCA
ITGB5.1	71	CC
JAG1.1	69	TGGCTTACACTGGCAATGGTAGTTTCTGTGGTTGGCTGGGAAATCGAGTGCCGCATCTCACAGCTATGC GTCAAAATGGGGAGGGACTAGGGCAGTTTGGATAGCTCAACAAGATACAATCTCACTCTGTGGTGGTCC
K-ras.10	71	TG
KCNJ15.1	67	GGACGTTCTACCTGCCTTGAAGAAGACACCTGACCTGCGGAGTGAGTGACCAGTGTTTCCAGAGCCT
KDR.6	68	GAGGACGAAGGCCTCTACACCTGCCAGGCATGCAGTGTTCTTGGCTGTGCAAAAGTGGAGGCATTTTT CGGACTTTGGGTGCGACTTGACGAGCGGTGGTTCGACAAGTGGCCTTGCGGGCCGGATCGTCCCAGTG
Ki-67.2	80	GAAGAGTTGTAA
KIAA1303 raptor.1	66	ACTACAGCGGGAGCAGGAGCTGGAGGTAGCTGCAATCAACCCAAATCACCTCTTGCTCAGATGCC
KIF1A.1	66	CTCCTACTGGTCGCACACCTCACCTGAGGACATCAACTACGCGTCGCAGAAGCAGGTGTACCGGGA GTCCCCGGGATGGATGTTTTGCCAAGTCATTGTTGGATAAGCGAGATGGTAGTACAATTGTCAGACAGC
Kitlng.4	79	TTGACTGATC GAGGTCCTGTCTAAACCCTGTGTCCCTGAGGGATCTGTCTCACTGGCATCTTGTTGAGGGCCTTGACAA
KL.1	72	TAG
KLK3.1	66	CCAAGCTTACCACCTGCACCCGGAGAGCTGTGTCAACCATGTGGGTCCCGTTGTCTTCCTCACCT
KLRK1.2	70	TGAGAGCCAGGCTTCTTGATGTCTCAAAATGCCAGCCTTCTGAAAGTATACAGCAAAGAGGACCAGGAT

Table B

Gene	Target Sequence Length	Amplicon Sequence
KRT19.3	77	TGAGCGGCAGAAATCAGGAGTACCAGCGGCTCATGGACATCAAGTCGCGGCTGGAGCAGGAGATTGCCA CCTACCGCA
KRT5.3	69	TCAGTGGAGAAGGAGTTGGACCAGTCAACATCTCTGTTGTCACAAGCAGTGTTCCTCTGGATATGGCA TTCAGAGATGAACCGGGCCATCCAGAGGCTGCAGGCTGAGATCGACAACATCAAGAACCAGCGTGCCAA
KRT7.1	71	GT
L1CAM.1	66	CTTGCTGGCCAATGCCTACATCTACGTTGTCCAGCTGCCAGCCAAGATCCTGACTGCGGACAATCA CAGATGAGGCACATGGAGACCCAGGCCAAGGACCTGAGGAATCAGTTGCTCAACTACCGTTCTGCCATT
LAMA3.1	73	TCAA
LAMA4.1	67	GATGCACTGCGGTTAGCAGCGCTCTCCATCGAGGAAGGCCAAATCCGGGGTGCTGAGCGTATCCTCTG
LAMB1.1	66	CAAGGAGACTGGGAGGTGTCTCAAGTGCCTGTACCACACGGAAGGGGAACACTGTCAGTTCTGCCG
LAMB3.1	67	ACTGACCAAGCCTGAGACCTACTGCACCCAGTATGGCGAGTGGCAGATGAAATGCTGCAAGTGTGAC ACTCAAGCGGAAATTGAAGCAGATAGGTCTTATCAGCACAGTCTCCGCCTCCTGGATTGAGTGTCTCGG
LAMC2.2	80	CTTCAGGGAGT
LAPTM5.1	66	TGCTGGACTTCTGCCTGAGCATCCTGACCCTCTGCAGCTCCTACATGGAAGTGCCACCTATCTCA
LDB1.2	67	AACACCCAGTTTGACGCAGCCAACGGCATTGACGACGAGGACAGCTTTAACAACCTCCCCTGCACTGG
LDB2.1	66	ATCACGGTGGACTGCGACCAGTGTACCATGGTCACCCAGCACGGAAGCCCATGTTTACCAAGGTA AGGCTACACATCCTGGGCTATTGGACTCTCTGTAGCAGATTTGGCAGAGAGTATAATGAAGAATCTTAGG
LDHA.2	74	CGGG
LGALS1.1	72	GGGTGGAGTCTTCTGACAGCTGGTGCGCCTGCCCCGGGAACATCCTCCTGGACTCAATCATGGCTTGTG GTCT
LGALS3.1	69	AGCGGAAAATGGCAGACAATTTTCGCTCCATGATGCGTTATCTGGGTCTGGAAACCCAAACCCTCAAG
LGALS9.1	67	AGTACTTCCACCGCGTGCCCTTCCACCGTGTGGACACCATCTCCGTCAATGGCTCTGTGCAGCTGTC
LIMK1.1	67	GCTTCAGGTGTTGTGACTGCAGTGCCTCCCTGTGCGACCAGTACTATGAGAAGGATGGGCAGCTCTT
LMNB1.1	66	TGCAAACGCTGGTGTACAGCCAGCCCCCAACTGACCTCATCTGGAAGAACCAGAACTCGTGGGG GGCTGCCAGCAGAACATCGGGGACCGCTACTTCCTGAAGGCCATCGACCAGTACTGGCACGAGGACTG
LMO2.1	74	CCTGAG
LOX.1	66	CCAATGGGAGAACAAACGGGCAGGTGTTGAGCTTGCTGAGCCTGGGCTCACAGTACCAGCCTCAGCG
LRP2.1	66	GGCTGTAGACTGGGTTTCCAGAAAGCTCTACTGGTTGGATGCCCGCCTGGATGGCCTCTTTGTCTC

Table B

Gene	Target Sequence Length	Amplicon Sequence
LRRC2.1	71	CCAGTGTCCCAATCTGTGTCCTGCGGATGTCTGAATTTGCAGTGGTTGGATATCAGCAGCAATAACCTGA CC
LTF.1	68	AACGGAAGCCTGTGACTGAGGCTAGAAGCTGCCATCTTGCCATGGCCCCGAATCATGCCGTGGTGTCT TTGCTGCAAGATAACATCGCTGATGCTGTAGCTTGTGCAAAGAGGGTGTCCGTGATCCACAAGGCATTA
LYZ.1	80	GAGCATGGGT GCTGCCTTTGGTAAGAACATGTCGTCCATCTTGCCATTACGCGGCCAGTTGTGAAGAGACTGCTGGGA
MADH2.1	70	T GGACATTACTGGCCTGTTCAATGAGCTTGCAATCCAGCCTCCCATTTCCAATCATCCTGCTCCTGAGT
MADH4.1	76	ATTGGT
MAL.1	66	GTTGGGAGCTTGCTGTGTCTAACCTCCAAGTGTGTGCTGTCTGCTAGGGTCACCTCCTGTTTGTG
MAL2.1	67	CCTTCGTCTGCCTGGAGATTCTGTTCCGGGGTCTTGTCTGGATTTTGGTTGCCTCCTCCAATGTTCC GCCTTTCTTACCCAGAAGCAGAAGGTGGGAGAACTGAAGGATGACGACTTTGAGAAGATCAGTGAGCTG
MAP2K1.1	76	GGGGCTG
MAP2K3.1	67	GCCCTCCAATGTCCTTATCAACAAGGAGGGCCATGTGAAGATGTGTGACTTTGGCATCAGTGGCTAC GCCGGTCAGGCACACAAGGGGCCCTTGGAGCGTGGACTGGTTGGTTTTGCCATTTTGTGTGTGTATGC
MAP4.1	72	TGC
MARCKS.1	67	CCCCTCTTGGATCTGTTGAGTTTCTTTGTTGAAGAAGCCAGCATGGGTGCCAGTTCTCCAAGACCG CAGATGGCCACTTTGAGAACATTTTAGCTGACAACAGTGTGAACGACCAGACCAAATCCTTGTGGTTAA
Maspin.2	77	TGCTGCC
MCAM.1	66	CGAGTTCCAGTGGCTGAGAGAAGAGACAGGCCAGGTGCTGGAAAGGGGGCCTGTGCTTCAGTTGCA GACTTTTGGCCGCTACCTTTTATTCCGGCGTGACAACAATGAGCTGTTGCTCTTCATACTGAAGCAGTTA
MCM2.2	75	GTGGC GGAGAACAAATCCCCTTGAGACAGAATATGGCCTTTCTGTCTACAAGGATCACCAGACCATCACCATCCAG
MCM3.3	75	GAGAT TGATGGTCCTATGTGTACATTCATCACAGTTTTCATACCAACACAGGCTTCAGCACTTCCTTTGGTGTG
MCM6.3	82	TTTCCTGTCCCA CGCTCAGCCAGATGCAATCAATGCCCCAGTCACCTGCTGTTATAACTTCACCAATAGGAAGATCTCAGTG
MCP1.1	71	C
MDH2.1	63	CCAACACCTTTGTTGCAGAGCTGAAGGGTTTGGATCCAGCTCGAGTCAACGTCCCTGTCATTG
MDK.1	66	GGAGCCGACTGCAAGTACAAGTTTGAGAACTGGGGTGCGTGTGATGGGGGCACAGGCACCAAAGTC
MDM2.1	68	CTACAGGGACGCCATCGAATCCGGATCTTGATGCTGGTGTAAAGTGAACATTCAGGTGATTGGTTGGAT

Table B

Gene	Target Sequence Length	Amplicon Sequence
MGMT.1	69	GTGAAATGAAACGCACCACACTGGACAGCCCTTTGGGGAAGCTGGAGCTGTCTGGTTGTGAGCAGGGT C
mGST1.2	79	ACGGATCTACCACACCATTGCATATTTGACACCCCTTCCCCAGCCAAATAGAGCTTTGAGTTTTTTTGTTG GATATGGA
MICA.1	68	ATGGTGAATGTCACCCGCAGCGAGGCCTCAGAGGGGCAACATTACCGTGACATGCAGGGCTTCTGGCTT
MIF.2	66	CCGGACAGGGTCTACATCAACTATTACGACATGAACGCGGCCAATGTGGGCTGGAACAACCTCCACC GGGAGATCATCGGGACAACCTCTCCTTTTGATGGACCTGGAGGAAATCTTGCTCATGCTTTTCAACCAGG
MMP1.1	72	CCC
MMP10.1	66	TGGAGGTGACAGGGAAGCTAGACACTGACACTCTGGAGGTGATGCGCAAGCCCAGGTGTGGAGTTC
MMP14.1	66	GCTGTGGAGCTCTCAGGAAGGGCCCTGAGGAAGGCACACTTGCTCCTGTTGGTCCCTGTCTTGCT CCATGATGGAGAGGCAGACATCATGATCAACTTTGGCCGCTGGGAGCATGGCGATGGATAACCCCTTTGA
MMP2.2	86	CGGTAAGGACGGACTCC
MMP7.1	79	GGATGGTAGCAGTCTAGGGATTAACCTCCTGTATGCTGCAACTCATGAACCTGGCCATTCTTTGGGTATG GGACATTCC
MMP9.1	67	GAGAACCAATCTCACCGACAGGCAGCTGGCAGAGGAATACCTGTACCGCTATGGTTACACTCGGGTG TCATGGTGCCCGTCAATGCTGTGATGGCGATGAAGACCAAGACGTATCAGGTGGCCACATGAAGAGCA
MRP1.1	79	AAGACAATCG
MRP2.3	65	AGGGGATGACTTGGACACATCTGCCATTGACATGACTGCAATTTTGACAAAGCCATGCAGTTTT TCATCCTGGCGATCTACTTCCTCTGGCAGAACCTAGGTCCCTCTGTCCTGGCTGGAGTCGCTTTCATGG
MRP3.1	91	TCTTGCTGATTCCACTCAACGG
MRP4.2	66	AGCGCCTGGAATCTACAACCTCGGAGTCCAGTGTTTTCCCACTTGTCATCTTCTCTCCAGGGGCTCT GATGCAGAATTGAGGCAGACTTTACAAGAAGATTTACTTCGTCGATTCCCAGATCTTAACCGACTTGCCA
MSH2.3	73	AGA
MSH3.2	82	TGATTACCATCATGGCTCAGATTGGCTCCTATGTTCTGCAGAAGAAGCGACAATTGGGATTGTGGATGG CATTTTCAACAG
MSH6.3	68	TCTATTGGGGGATTGGTAGGAACCGTTACCAGCTGGAAATTCCTGAGAATTTCACCACTCGCAATTTG
MT1B.1	66	GTGGGCTGTGCCAAGTGTGCCAGGGCTGTGTCTGCAAAGGCTCATCAGAGAAGTGCCGCTGCTGT GTGCACCCACTGCCTCTTCCCTTCTCGCTTGGGAACTCTAGTCTCGCCTCGGGTTGCAATGGACCCCAA
MT1G.1	74	CTGCT
MT1H.1	74	CGTGTTCACCTGCCTCTTCTCTTCTCGCTTGGGAACTCCAGTCTCACCTCGGCTTGCAATGGACCCCAAC

Table B

Gene	Target Sequence Length	Amplicon Sequence
MT1X.1	80	CTCCTGCAAATGCAAAGAGTGCAAATGCACCTCCTGCAAGAAGAGCTGCTGCTCCTGCTGCCCTGTGGGCTGTGCCAAGT
MUC1.2	71	GGCCAGGATCTGTGGTGGTACAATTGACTCTGGCCTTCCGAGAAGGTACCATCAATGTCCACGACGTGGAG
MVP.1	75	ACGAGAACGAGGGCATCTATGTGCAGGATGTCAAGACCGGAAAGGTGCGCGCTGTGATTGGAAGCACCACATGC
MX1.1	78	GAAGGAATGGGAATCAGTCATGAGCTAATCACCTGGAGATCAGCTCCCGAGATGTCCCGGATCTGACTCTAATAGAC
MYBL2.1	74	GCCGAGATCGCCAAGATGTTGCCAGGGAGGACAGACAATGCTGTGAAGAATCACTGGAACTCTACCATCGCCGAGATCGCCAAGATGTTGCCAGGGAGGACAGACAATGCTGTGAAGAATCACTGGAACTCTACCATC
MYH11.1	85	CGGTACTTCTCAGGGCTAATATATACGTACTCTGGCCTCTTCTGCGTGGTGGTCAACCCCTATAAACACCTGCCCCATCTACTCGG
MYRIP.2	69	CCTTCACCTTCCTCGTCAACACCAAGCGCCAGTGTGGAGATTGCAAATTCAATGTCTGCAAGAGCTGCTGCATCTACTTGCCAGAACCAAATTACTTACTTCCAAGTTCTGGCTGCTTGCAGGTGGAAGTCCAGCTGC
NBN.1	76	AAGGGA
NCF1.1	66	GACACCTTCATCCGTACATCGCCCTGCTGGGCTTTGAGAAGCGCTTCGTACCCAGCCAGCACTATCTGAACCCCTCTCCTGGTCACCGAGAATCAGTCCCCGTGGAGTTCCCCCTCCACCTCGCCATCGTTTCC
NFAT5.1	70	T
NFATC2.1	72	AAAG
NFKBp50.3	73	AGCT
NFKBp65.3	68	CTGCCGGGATGGCTTCTATGAGGCTGAGCTCTGCCCGGACCGCTGCATCCACAGTTTCCAGAACCTGGCCCTGCCATACCAGCTCACCTGCCCCTGTGACTGCTTGTAAGCTAAGGTAGAGCTACAGTGTGAATGT
NFX1.1	74	GGACG
NME2.1	66	ATGCTTGGGGAGACCAATCCAGCAGATTCAAAGCCAGGCACCATTCGTGGGGACTTCTGCATTGAG
NNMT.1	67	CCTAGGGCAGGGATGGAGAGAGAGTCTGGGCATGAGGAGAGGGTCTCGGGATGTTTGGCTGGACTAGCAGCCTTGGGAAGTGAGACTAGAAGAGGGGAGCAGAAAGGGACCTTGAGTAGACAAAGGCCACACACA
NOL3.1	72	TCAT
NOS2A.3	67	GGGTCCATTATGACTCCCAAAAGTTTGACCAGAGGACCCAGGGACAAGCCTACCCCTCCAGATGAGC

Table B

Gene	Target Sequence Length	Amplicon Sequence
NOS3.1	68	ATCTCCGCCTCGCTCATGGGCACGGTGATGGCGAAGCGAGTGAAGGCGACAATCCTGTATGGCTCCGA CGGGTCCACCAGTTTGAATGGTCAATGCGAGTGGCTGTCCCGGCTGCAGAGCGGCATGGTGCCGAACC
NOTCH1.1	76	AATACAAC CACTTCCTGCTGGGATTATATCAACAACCAGTGTGATGAGCTGTGCAACACGGTCGAGTGCCTGTTTG
NOTCH2.1	75	ACAAC
NOTCH3.1	67	TGTGGACGAGTGTGCTGGCCCCGCACCCTGTGGCCCTCATGGTATCTGCACCAACCTGGCAGGGAGT GGCTGTGGCTGAGGCTGTAGCATCTCTGCTGGAGGTGAGACACTCTGGGAACTGATTTGACCTCGAATG
NPD009 (ABAT official).3	73	CTCC AATGTTGTCCAGGTTCTATTGCCAAGAATGTGTTGTCCAAATGCCTGTTTAGTTTTAAAGATGGAACTC
NPM1.2	84	CACCCTTTGCTTG
NPPB.1	66	GACACCTGCTTCTGATTCCACAAGGGGCTTTTTCTCAACCCTGTGGCCGCCTTTGAAGTGA
NPR1.1	66	ACATCTGCAGCTCCCCTGATGCCTTCAGAACCCTCATGCTCCTGGCCCTGGAAGCTGGCTTGTGTG
NPY1R.1	70	GGATCTTCCCCACTCTGCTCCCTTCCATTCCCACCCTTCCTTCTTTAATAAGCAGGAGCGAAAAAGACAA CGAGACTCTCCTCATAGTGAAAGGTATGTGTCAGCCATGACCACCCCGGCTCGTATGTCACCTGTAGATT
NRG1.3	83	TCCACACGCCAAG ACTGGTTTCCACTCCTGCTTCAGAAGAAGAAATTCCACGGGTACTTCAAGTTCAGGGTCAGGACACCAT
NUDT1.1	77	CCTGGAC ACCAAGGTGGCTGACTGCATCTGCCTGATGGCCCTAGACAAGCCCCAGGCTGTGCCCGTGGATGTCCA
OGG1.1	71	TAT CAACCGAAGTTTTCACTCCAGTTGTCCCCACAGTAGACACATATGATGGCCGAGGTGATAGTGTGGTTTA
OPN, osteopontin.3	80	TGGACTGAGG
p21.3	65	TGGAGACTCTCAGGGTCGAAAACGGCGGCAGACCAGCATGACAGATTTCTACCACTCCAAACGCC
p27.3	66	CGGTGGACCACGAAGAGTTAACCCGGGACTTGGAGAAGCACTGCAGAGACATGGAAGAGGCGAGCC
P53.2	68	CTTTGAACCCTTGCTTGCAATAGGTGTGCGTCAGAAGCACCCAGGACTTCCATTTGCTTTGCCCGGG TGGCTGATTCCATTAACAGTGAAATTGGAATCCTTTGCAGTGCCCTCCAGAAAATAAAGTAAAGCCATGG
PAH.1	80	ACAGAATGTG CCGCAACGTGGTTTTCTCACCTATGGGGTGGCCTCGGTGTTGGCCATGCTCCAGCTGACAACAGGAG
PAI1.3	81	GAGAAACCCAGCA GAGCTGTGGGTTGTTATGGAATACTTGGCTGGAGGCTCCTTGACAGATGTGGTGACAGAACTTGCATG
Pak1.2	70	G

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Gene	Target Sequence Length	Amplicon Sequence
PARD6A.1	66	GATCCTCGAGGTCAATGGCATTGAAGTAGCCGGGAAGACCTTGGACCAAGTGACGGACATGATGGT GCAAAGCCTTTCCAGAAAAATAAAAATGGTTGAAAAGGCAATTCTGCTACCAATGACTGTTTAAGCCCAG
PBOV1.1	72	CC
PCCA.1	68	GGTGAAATCTGTGCACTGTCAAGCTGGAGACACAGTTGGAGAAGGGGATCTGCTCGTGGAGCTGGAAT
PCK1.1	66	CTTAGCATGGCCCAGCACCCAGCAGCCAAACTGCCCAAGATCTTCCATGTCAACTGGTTCCGGAAG GAAGGTGTTGGAGGCACTCAAGGACCTCATCAACGAGGCCTGCTGGGATATTAGCTCCAGCGGTGTAA
PCNA.2	71	CC
PCSK6.1	67	ACCTTGAGTAGCAGAGGCCCTCACACCTTCCTCAGAATGGACCCCCAGGTGAAATGGCTCCAGCAAC GACAACGCCACCTTCACCTGCAGCTTCTCCAACACATCGGAGAGCTTCGTGCTAAACTGGTACCGCATG
PDCD1.1	73	AGCC GCTTCGTCTTGCTGTGAGAGAGCGAGATCATGACTTAGAGAGACTGCGCGATGTCCTCTCCTCCAATGA
PDE4DIP.1	73	AGCT
PDGFA.3	67	TTGTTGGTGTGCCCTGGTGCCGTGGTGGCGGTCACTCCCTCTGCTGCCAGTGTTTGGACAGAACCCA
PDGFB.3	62	ACTGAAGGAGACCCTTGGAGCCTAGGGGCATCGGCAGGAGAGTGTGTGGGCAGGGTTATTTA AGTTACTAAAAAATACCACGAGGTCCTTCAGTTGAGACCAAAGACCGGTGTCAGGGGATTGCACAAATCA
PDGFC.3	79	CTCACCGAC
PDGFD.2	74	TATCGAGGCAGGTCATACCATGACCGGAAGTCAAAAGTTGACCTGGATAGGCTCAATGATGATGCCAAG CGTTA GGGAGTTTCCAAGAGATGGACTAGTGCTTGGTCGGGTCTTGGGGTCTGGAGCGTTTGGGAAGGTGGTT
PDGFRa.2	72	GAAG
PDGFRb.3	66	CCAGCTCTCCTTCCAGCTACAGATCAATGTCCCTGTCCGAGTGCTGGAGCTAAGTGAGAGCCACCC AATGACCTCCACCTTCAACCCCCGAGAATGTAAACTGTCCAAGCAAGAAGGGCAAACTATGGCTTCTTC
PDZK1.1	75	CTGCG
PDZK3.1	68	GAGCTGAGAGCCTTGAGCATGCCTGACCTTGACAAGCTCTGCAGCGAGGATTACTCAGCAGGGCCGAG GCAGTGCCTGTGTGTGAAGACCACCTCCCAGGTCCGTCCCAGGCACATCACCAGCCTGGAGGTGATCA
PF4.1	73	AGGCC
PFKP.1	68	AGCTGATGCCGCATACATTTTCGAAGAGCCCTTCGACATCAGGGATCTGCAGTCCAACGTGGAGCACC TCTATACGTGATGGTGACTGCACAATGGACATCCGGACAAAGAGTCAAGGTGGGGAGCCAACATACAA
PFN2.1	82	TGTGGCTGTGGC

Table B

Gene	Target Sequence Length	Amplicon Sequence
PGF.1	71	GTGGTTTTCCCTCGGAGCCCCCTGGCTCGGGACGTCTGAGAAGATGCCGGTCATGAGGCTGTTCCCTT GCT
PI3K.2	98	TGCTACCTGGACAGCCCGTTGGTGCGCTTCCTCCTGAAACGAGCTGTGTCTGACTTGAGAGTGA CTAC TACTTCTTCTGGTTACTGAAGGACGGCCT
PI3KC2A.1	83	ATACCAATCACCGCACAAACCCAGGCTATTTGTTAAGTCCAGTCACAGCACAAAGAAACATATGCGGAGA AAATGCTAGTGTG
PIK3CA.1	67	GTGATTGAAGAGCATGCCAATTGGTCTGTATCCCGAGAAGCAGGATTTAGCTATTCCCACGCAGGAC
PLA2G4C.1	68	CCCTTTCCCCAAGTAGAAGAGGCTGAGCTGGATTTGTGGTCCAAGGCCCGCCAGCTGCTACATCCT
PLAT.1	67	GATTTGCTGGGAAGTGCTGTGAAATAGATACCAGGGCCACGTGCTACGAGGACCAGGGCATCAGCTA CCCATGGATGCTCCTCTGAAGAGACTTTCCTCATTGACTGCCGAGGCCCATGAATCAATGTCTGGTAG
PLAUR.3	76	CCACCGG
PLG.1	77	GGCAAAATTTCCAAGACCATGTCTGGACTGGAATGCCAGGCCTGGGACTCTCAGAGCCACACGCTCAT GGATACAT
PLN.1	84	TGATGCTTCTCTGAAGTTCTGCTACAACCTCTAGATCTGCAGCTTGCCACATCAGCTTAAAATCTGTCATC CCATGCAGACAGG
PLOD2.1	84	CAGGGAGGTGGTTGCAAATTTCTAAGGTACAATTGCTCTATTGAGTCACCACGAAAAGGCTGGAGCTTCA TGCATCCTGGGAGA
PLP1.1	66	AGAACAGACTGGCCTGAGGAGCAGCAGTTGCTGGTGGCTAATGGTGTAACCTGAGATGGCCCTCTG
PMP22.1	66	CCATCTACACGGTGAGGCACCCGGAGTGGCATCTCAACTCGGATTACTCCTACGGTTTCGCCTACA ACAAGCACCATCCCAGTGATGTTCTGGCAGGATTTGCTCAAGGAGCCCTGGTGGCCTGCTGCATAGTTT
PPAP2B.1	77	TCTTCGTG
PPARG.3	72	TGACTTTATGGAGCCCAAGTTTGAGTTTGCTGTGAAGTTCAATGCACTGGAATTAGATGACAGCGACTTG GC
PPP1R3C.1	82	TTCCTTCCCTCTCAATCCACTAGCTTTCATGTTGGGCAAGGAAAAGTTGAGGAAGGATGGCTGATGGTGA TGGAAGCTGTG
PPP2CA.1	78	GCAATCATGGAAGTTGACGATACTCTAAAATACTCTTCTTGACAGTTTGACCCAGCACCTCGTAGAGGCG AGCCACAT
PRCC.1	67	GAGGAAGAGGAGGCGGTGGCTCCTACATCTGGGCCCGCTTTAGGGGGCTTGTTTCGCTTCTCTCCCTG CAAGCAATGCGTCATCAATGTCCCCAGCCTCTGCGGAATGGATCACACTGAGAAGAGGGGGCGGATTTA
PRKCA.1	70	C
PRKCB1.1	67	GACCCAGCTCCACTCCTGCTTCCAGACCATGGACCGCCTGTACTTTGTGATGGAGTACGTGAATGGG
PRKCD.2	68	CTGACACTTGCCGCAGAGAATCCCTTCTCACCCACCTCATCTGCACCTTCCAGACCAAGGACCACCT

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Gene	Target Sequence Length	Amplicon Sequence
PRKCH.1	68	CTCCACCTATGAGCGTCTGTCTCTGTGGGCTTGGGATGTTAACAGGAGCCAAAAGGAGGGAAAGTGTG
PRO2000.3	79	ATTGGA AAAACCTCGTCACCAGAGAAAGGCCAACATATTTATAGTGGCCCAGCTTCTCCTGCAAGACCA AGATAACCGA
PROM1.1	74	CTATGACAGGCATGCCACCCCGACCACCCGAGGCTGTGTCTCCAACACCCGAGGCGTCTTCCTCATGGT TGGAG
PROM2.1	67	CTTCAGCGCATCCACTACCCCGACTTCCTCGTTCAGATCCAGAGGCCCGTGGTGAAGACCAGCATGG
PRPS2.1	69	CACTGCACCAAGATTCAGGTCATTGACATTTCCATGATCTTGGCCGAAGCAATCCGAAGGACACACAAT
PRSS8.1	68	GTACACTCTGGCCTCCAGCTATGCCTCCTGGATCCAAAGCAAGGTGACAGAACTCCAGCCTCGTGTGG
PSMA7.1	67	GCCAAACTGCAGGATGAAAGAACAGTGCGGAAGATCTGTGCTTTGGATGACAACGTCTGCATGGCCT
PSMB8.1	66	CAGTGGCTATCGGCCTAATCTTAGCCCTGAAGAGGCCTATGACCTTGGCCGCAGGGCTATTGCTTA
PSMB9.1	66	GGGGTGTCTATCTACCTGGTCACTATTACAGCTGCCGGTGTGGACCATCGAGTCATCTTGGGCAATG TGGCTAAGTGAAGATGACAATCATGTTGCAGCAATTCACTGTAAAGCTGGAAAGGGACGAAGTGGTGTA
PTEN.2	81	TGATATGTGCA
PTGIS.1	66	CCACACTGGCATCTCCCTGACCTTCTCCAGGGACAGAAGCAGGAGTAAGTTTCTCATCCCATGGGC CGAGGTACAAGCTGAGATCAAGAAATCTTGGAGCCGCTGGACACTGGCACTGGACTTCAAGCGAAAGGC
PTHR1.1	73	ACGC
PTK2.1	68	GACCGGTGCAATGATAAGGTGTACGAGAATGTGACGGGCCTGGTGAAAGCTGTCATCGAGATGTCCAG CAAGCCCAGCCGACCTAAGTACAGACCCCTCCGCAAACCAACCTCCTGGCTCCAAAGCTGCAGTTCCA
PTK2B.1	74	GGTTC
PTN.1	67	CCTTCCAGTCCAAAAATCCCGCCAAGAGAGCCCCAGAGCAGAGGAAAAATCCAAAGTGGAGAGAGGGG CTCCAGCTAGCACTAAGCAACATCTCGCTGTGGACGCTGTAAATTACTGAGAAATGTGAAACGTGCAAT
PTPNS1.1	77	CTTGAAA
PTPRB.1	68	GATATGCGGTGAGGAACAGCTTGATGCACACAGACTCATCCGCCACTTTCACTATACGGTGTGGCCAG TGGCCGTCAATGGAAGAGGGGCACTCGGGCTTTTGGAGATGTTGTTGTAAGATCAACCAGCACAAAAGA
PTPRC.1	74	TGTCC
PTPRG.1	71	GGACAGCGACAAAGACTTGAAAGCCACCATTAGCCATGTCTCACCCGATAGCCTTTACCTGTTCCGAGTC C

Table B

Gene	Target Sequence Length	Amplicon Sequence
PTTG1.2	74	GGCTACTCTGATCTATGTTGATAAGGAAAATGGAGAACCAGGCACCCGTGTGGTTGCTAAGGATGGGCT GAAGC
PVALB.1	81	AAACCAAGATGCTGATGGCTGCTGGAGACAAAGATGGGGACGGCAAATTTGGGGTTGACGAATTCTCCA CTCTGGTGGCTG
PXDN.1	67	GCTGCTCAAGCTGAACCCGCACTGGGACGGCGACACCATCTACTATGAGACCAGGAAGATCGTGGGT
RAC1.3	66	TGTTGTAAATGTCTCAGCCCTCGTTCTTGGTCCTGTCCCTTGGAACCTTTGTACGCTTTGCTCAA
RAD51.1	66	AGACTACTCGGGTCGAGGTGAGCTTTCAGCCAGGCAGATGCACTTGGCCAGGTTTCTGCGGATGCT CGTCGTATGCGAGAGTCTGTTTCCAGGATGCCTGTTAGTTCTCAGCACAGATATTCTACACCTCACGCCT
RAF1.3	73	TCA
RALBP1.1	84	GGTGTGAGATATAAATGTGCAAATGCCTTCTTGCTGTCCTGTGCGTCTCAGTACGTTCACTTTATAGCTG CTGGCAATATCGAA
RARB.2	78	TGCCTGGACATCCTGATTCTTAGAATTTGCACCAGGTATACCCAGAACAAGACACCATGACTTTCTCAG ACGGCCTT
RASSF1.1	75	AGGGCACGTGAAGTCATTGAGGCCCTGCTGCGAAAGTTCTTGGTGGTGGATGACCCCCGCAAGTTTGC ACTCTTT
RB1.1	77	CGAAGCCCTTACAAGTTTCCTAGTTCACCCTTACGGATTCTGAGGGAACATCTATATTTACCCCTGA AGAGTCC
RBM35A.1	66	TGGTTTTGAATCACCAGGGCCGCCATCAGGAGATGCCTTTATCCAGATGAAGTCTGCGGACAGAG TGCTAACTCCTGCACAGCCCCGTCCTCTTCTTTCTGCTAGCCTGGCTAAATCTGCTCATTATTTACAGAG
REG4.1	83	GGGAAACCTAGCA
RET.1	71	GCCTGTGCAGTTCTTGTGCCCCAACATCAGCGTGGCCTACAGGCTCCTGGAGGGTGAGGGTCTGCCCT TCC
RGS1.1	84	TGCCCTGTAAAGCAGAAGAGATATATAAAGCATTTGTGCATTGATGCTGCTAAACAAATCAATATTGAC TTCCGCACTCGAG
RGS5.1	79	TTCAAACGGAGGCTCCTAAAGAGGTGAATATTGACCACTTCACTAAGGACATCACAATGAAGAACCTGGT GGAACCTTC
RHEB.2	78	GATGATTGAGAACAGCCTTGCCTGTCACTGTCTAGAACACCCTGGAGTTTAGTGTTCTGTGTCAGAGTC TTGGGAGC
RhoB.1	67	AAGCATGAACAGGACTTGACCATCTTTCCAACCCCTGGGGAAGACATTTGCAACTGACTTGGGGAGG
rhoC.1	68	CCCGTTTCGGTCTGAGGAAGGCCGGGACATGGCGAACCGGATCAGTGCCTTTGGCTACCTTGAGTGCTC
RIPK1.1	67	AGTACCTTCAAGCCGGTCAAATTCAGCCACAGAACAGCCTGGTTCACTGCACAGTTCCAGGGACTT
RND3.1	66	TCGGAATTGGACTTGGGAGGCGCGGTGAGGAGTCAGGCTTAAACTTGTTGGAGGGGAGTAACCAG

Table B

Gene	Target Sequence Length	Amplicon Sequence
ROCK1.1	73	TGTGCACATAGGAATGAGCTTCAGATGCAGTTGGCCAGCAAAGAGAGTGATATTGAGCAATTGCGTGCT
ROCK2.1	66	AAAC GATCCGAGACCCTCGCTCCCCCATCAACGTGGAGAGCTTGCTGGATGGCTTAAATTCCTTGGTCCT
RPLP1.1	68	CAAGGTGCTCGGTCCTTCCGAGGAAGCTAAGGCTGCGTTGGGGTGAGGCCCTCACTTCATCCGGCGAC
RPS23.1	67	GTTCTGGTTGCTGGATTTGGTCGCAAAGGTCATGCTGTTGGTGATATTCCTGGAGTCCGCTTTAAGG
RPS27A.1	74	CTTACGGGGAAGACCATCACCTCGAGGTTGAACCTCGGATACGATAGAAAATGTAAAGGCCAAGATC CAGGA
RPS6KAI.1	70	GCTCATGGAGCTAGTGCCTCTGGACCCGGAGAATGGACAGACCTCAGGGGAAGAAGCTGGACTTCAGC CG
RPS6KB1.3	81	GCTCATTATGAAAAACATCCCAAACCTTTAAATGCGAAATTATTGGTTGGTGTGAAGAAAGCCAGACAACT TCTGTTTCTT
RRM1.2	66	GGGCTACTGGCAGCTACATTGCTGGGACTAATGGCAATTCCAATGGCCTTGTACCGATGCTGAGAG
RRM2.1	71	CAGCGGGATTAAACAGTCCTTTAACCAGCACAGCCAGTTAAAAGATGCAGCCTCACTGCTTCAACGCAGA T
RUNX1.1	70	AACAGAGACATTGCCAACCATATTGGATCTGCTTGCTGTCCAAACCAGCAAACCTTTCCTGGGCAAATCAC TGGACAAGGTGATGAAGGAGCTAGACGAGAATGGAGACGGGGAGGTGGACTTCCAGGAGTATGTGGTG
S100A1.1	70	CT ACACCAAAAATGCCATCTCAAATGGAACACGCCATGGAAACCATGATGTTTACATTTACAAATTGCTGG
S100A10.1	77	GGATAAA TGGCTGTGCTGGTCACTACCTTCCACAAGTACTCCTGCCAAGAGGGCGACAAGTTCAAGCTGAGTAAGG
S100A2.1	73	GGGA CTACAGCACAGATCAGCACCATGAAGCTTCTCACGGGCCTGGTTTTCTGCTCCTTGGTCCTGAGTGTCA
SAA2.2	72	GCA GCCTTCCTGGAGTACCCGAGTGCTCCCTATGCCTTTCCAAGCATTTCTACTTGGGGAATTGGGCCAC
SCN4B.1	67	ATCAACATCCTGTGCGAGGCTGCCAGAGACTCTGCCATCCCTGGAGGAGGACACGCTGGGCAACTC
SCNN1A.2	66	
SDHA.1	67	GCAGAACTGAAGATGGGAAGATTTATCAGCGTGCATTTGGTGGACAGAGCCTCAAGTTTGGAAAGGG
SDPR.1	66	ACCAGCACAAGATGGAGCAGCGACAGATCAGTTTGGAGGGCTCCGTGAAGGGCATCCAGAATGACC ACACTGGTCTGGCCTGCTACCTACCTGTGAAGCTCCCACTGAGTCCAACATTCCCTTGGTAGCTGGACT
SELE.1	71	TT
SELENBP1.1	67	GGTACCAGCCTCGACACAATGTCATGATCAGCACTGAGTGGGCAGCTCCCAATGTCTTACGAGATGG

Table B

Gene	Target Sequence Length	Amplicon Sequence
SELL.1	67	TGCAACTGTGATGTGGGGTACTATGGGCCCCAGTGTGAGTTTGTGATTCAGTGTGAGCCTTTGGAGG
SELPLG.1	83	TGGCCACTATCTTCTTCGTGTGCACTGTGGTGTGGCGGTCCGCCTCTCCCCGAAGGGCCACATGTACC CCGTGCGTAATTAC
SEMA3B.1	71	GCTCCAGGATGTGTTTCTGTTGTCCTCGCGGGACCACCGGACCCCGCTGCTCTATGCCGTCTTCTCCAC
SEMA3C.1	66	GT ATGGCCATTCTGTTCCAGATTCTACCCAACTGGGAAACGGAGGAGCCGAAGACAAGATGTGAGAC
SEMA3F.3	86	CGCGAGCCCTCATTATACACTGGGCAGCCTCCCCACAGCGCATCGAGGAATGCGTGTCTCTCAGGCAA GGATGTCAACGGCGAGTG
SEMA5B.1	67	CTCGAGGACAGCTCCAACATGAGCCTCTGGACCCAGAACATCACCGCCTGTCTGTGCGGAATGTGA
SERPINA5.1	66	CAGCATGGTAGTGGCAAAGAGAGGTCCAGAGTCCTGGCCCTTGATGCCAGCTCAGTGCCACAAAG GAGAGAGCCAGTCTGATCCAGAAGGCCAAGCTGGCAGAGCAGGCCGAACGCTATGAGGACATGGCAGC
SFN.1	70	CT TCCGCAAGACACCTCCTGGAGGGCCTCCTGCAGAAGGACAGGACAAAGCGGCTCGGGGCCAAGGATGA
SGK.1	73	CTTCA
SHANK3.1	68	CTGTGCCCTCTACAACCAGGAGAGCTGTGCTCGTGTCTCTTCCGTGGAGCTAACAGGGATGTCC CCAACACCTTCTTGGCTTCTGGGACCTGTGTTCTTGCTGAGCACCTCTCCGGTTTGGGTTGGGATAAC
SHC1.1	71	AG
SILV.1	66	CCGCATCTTCTGCTCTTGTCCCATTGGTGAGAATAGCCCCCTCCTCAGTGGGCAGCAGGTCTGAGT
SKIL.1	66	AGAGGCTGAATATGCAGGACAGTTGGCAGAACTGAGGCAGAGATTGGACCATGCTGAGGCCGATAG
SLC13A3.1	66	CTTGCCCTCCAACAAGGTCTGCCCCCAGTACTTCCTCGACACCAACTTCCTCTTCCTCAGTGGGCT
SLC16A3.1	68	ATGCGACCCACGTCTACATGTACGTGTTTCATCCTGGCGGGGGCCGAGGTGCTCACCTCCTCCCTGATT
SLC22A3.1	66	ATCGTCAGCGAGTTTGACCTTGTCTGTGTCAATGCGTGGATGCTGGACCTACCCAAGCCATCCTG
SLC22A6.1	68	TCCGCCACCTCTTCCTCTGCCTCTCCATGCTGTGTTTGCCACTAGCTTTGCATACTATGGGCTGGTC
SLC2A1.1	67	GCCTGAGTCTCCTGTGCCACATCCCAGGCTTCACCTGAATGGTTCCATGCCTGAGGGTGGAGACT
SLC34A1.1	66	GCTGAGACCCACTGACCTGCAGACCTCATAGTGGGTGCCAGGATGTTGTCTACGGAGAGAGGCT GCGCAGAGGCCAGTTAAAGTAGATCACCTCCTCGAACCCACTCCGGTTCCCCGCAACCCACAGCTCAGC
SLC7A5.2	70	T
SLC9A1.1	67	CTTCGAGATCTCCCTCTGGATCCTTCTGGCCTGCCTCATGAAGATAGGTTTCCATGTGATCCCCACT
SLIT2.2	67	TTTACCGATGCACCTGTCCATATGGTTTCAAGGGGCAGGACTGTGATGTCCCAATTTCATGCCTGCAT

Table B

Gene	Target Sequence Length	Amplicon Sequence
SNAI1.1	69	CCCAATCGGAAGCCTAACTACAGCGAGCTGCAGGACTCTAATCCAGAGTTTACCTTCCAGCAGCCCTAC
SNRK.1	71	GAGGAAAAGTCAGGGCCGGGGCTCCAGCTGCAGTAGTTCGGAGACCAGTGATGATGATTCTGAAAGCC
SOD1.1	70	GGC TGAAGAGAGGCATGTTGGAGACTTGGGCAATGTGACTGCTGACAAAGATGGTGTGGCCGATGTGTCTAT
SP3.1	69	TCAAGAGTCTCAGCAGCCAACCAGTCAAGCCCAAATTGTGCAAGGTATTACACCACAGACAATCCATGG
SPARC.1	73	TCTTCCCTGTACACTGGCAGTTCGGCCAGCTGGACCAGCACCCCATTGACGGGTACCTCTCCCACACCG
SPARCL1.1	67	AGCT GGCACAGTGCAAGTGATGACTACTTCATCCCAAGCCAGGCCTTTCTGGAGGCCGAGAGAGCTCAATC
SPAST.1	66	CCTGAGTTGTTTACAGGGCTTAGAGCTCCTGCCAGAGGGCTGTTACTCTTTGGTCCACCTGGGAAT
SPHK1.1	67	GGCAGCTTCCTTGAACCATTATGCTGGCTATGAGCAGGTCACCAATGAAGACCTCCTGACCAACTGC
SPRY1.1	77	CAGACCAGTCCCTGGTCATAGGTCTGAAAGGGCAATCCGGACCCAGCCCAAGCAACTGATTGTGGATGA
SQSTM1.1	69	CTTGAAGG GGACCCGTCTACAGGTGAACTCCAGTCCCTACAGATGCCAGAATCCGAAGGGCCAAGCTCTCTGGACCC
STAT1.3	81	GGGCTCAGCTTTCAGAAGTGCTGAGTTGGCAGTTTTCTTCTGTACCAAAGAGGTCTCAATGTGGACCA
STAT3.1	70	GCTGAACATGT TCACATGCCACTTTGGTGTTTTCATAATCTCCTGGGAGAGATTGACCAGCAGTATAGCCGCTTCCTGCAAG
STAT5A.1	77	GAGGCGCTCAACATGAAATTCAAGGCCGAAGTGACAGAGCAACCGGGGCTGACCAAGGAGAACCTCGT
STAT5B.2	74	GTTCTCTGGC CCAGTGGTGGTGATCGTTCATGGCAGCCAGGACAACAATGCGACGGCCACTGTTCTCTGGGACAATGCT
STC2.1	67	TTTGC AAGGAGGCCATCACCCACAGCGTGCAGGTTCAAGTGTGAGCAGAACTGGGGAAGCCTGTGCTCCATCT
STK11.1	66	GGAAGGATCCC GGAICTCGGAGACGCTGTGCAGGAGGGCCGTCAAGATCCTCAAGAAGAAGAAGTTGCGAAGGATCCC
STK15.2	69	CATCTTCCAGGAGGACCACTCTCTGTGGCACCCCTGGACTACCTGCCCCCTGAAATGATTGAAGGTCCGA
STK4.1	66	GAGCCATCTTCTGCAACTTTACCTCTTTCCCTCAGATGGGAGCCATGACTGGGTTGCACCTCAG
STMY3.3	90	CCTGGAGGCTGCAACATACCTCAATCCTGTCCCAGGCCGGATCCTCCTGAAGCCCTTTTCGCAGCACTG
		CTATCCTCCAAAGCCATTGTA

Table B

Gene	Target Sequence Length	Amplicon Sequence
SUCLG1.1	66	CCAAGCCTGTAGTGTCTTCATTGCTGGTTTAACTGCTCCTCCTGGGAGAAGAATGGGTTCATGCCG
SULT1C2.1	67	GGGACCCAAAGCATGAAATTCGGAAGGTGATGCAGTTCATGGGAAAGAAGGTGGATGAAACAGTGCT
SURV.2	80	TGTTTTGATTCCCGGGCTTACCAGGTGAGAAGTGAGGGAGGAAGAAGGCAGTGTCCTTTTGCTAGAGC
TACSTD2.1	80	ATCACCAACCGGAGAAAGTCGGGGAAGTACAAGAAGGTGGAGATCAAGGAACTGGGGGAGTTGAGAAA
TAGLN.1	73	GGAACCGAGCTT GATGGAGCAGGTGGCTCAGTTCCTGAAGGCGGCTGAGGACTATGGGGTCATCAAGACTGACATGTTCC
TAP1.1	72	AGACT GTATGCTGCTGAAAGTGGGAATCCTCTACATTGGTGGGCAGCTGGTGACCAGTGGGGCTGTAAGCAGT
TCF4.1	67	GGG
TCOF1.2	66	CACACCCTGGAATGGGAGACGCATCGAATCACATGGGACAGATGTAAAAGGGTCCAAGTTGCCACAT
TEK.1	76	AGCGAGGATGAGGACGTGATCCCCGCTACACAGTGCTTGACTCCTGGCATCAGAACCAATGTGGTG
TERT.1	85	ACTTCGGTGCTACTTAACAACCTACATCCCAGGGAGCAGTACGTGGTCCGAGCTAGAGTCAACACCAAG
TFAP2B.1	67	GCCCCAGG GACATGGAGAACAAGCTGTTTGCGGGGATTTCGGCGGGACGGGCTGCTCCTGCGTTTGGTGGATGATTT
TFAP2C.1	68	CTTGTTGGTGACACCTC
TFPI.1	69	CGTGTGACGTGCGAGAGACGCGATGGACGCGCCTTGCTCTTACTGTGCAGGTCCTGAGAGCGTGTGG
TGFA.2	83	CATGCCTCACCAGATGGACGAGGTGCAGAATGTGACGACCAGCACCTGTTGCTGCACGATCAGACAG
TGFb1.1	80	CCGAATGGTTTCCAGGTGGATAATTATGGAACCCAGCTCAATGCTGTGAATAACTCCCTGACTCCGCAA
TGFB2.2	75	GGTGTGCCACAGACCTTCCTACTTGGCCTGTAATCACCTGTGCAGCCTTTTGTGGGCCTTCAAACTCTG
TGFBI.1	67	TCAAGAACTCCGT CTGTATTTAAGGACACCCGTGCCCAAGCCCACCTGGGGCCCCATTAAAGATGGAGAGAGGACTGCGG
TGFBR1.1	67	ATCTCTGTGTCA ACCAGTCCCCCAGAAGACTATCCTGAGCCCGAGGAAGTCCCCCGGAGGTGATTTCCATCTACAACAGC
TGFBR2.3	66	ACCAGG GCTACGAGTGCTGTCCTGGATATGAAAAGGTCCCTGGGGAGAAGGGCTGTCCAGCAGCCCTACCACT
	67	GTCATCACCTGGCCTTGGTCCTGTGGAAGTGGCAGCTGTCATTGCTGGACCAGTGCTGCTTCGTCTGC
	66	AACACCAATGGGTTCCATCTTTCTGGGCTCCTGATTGCTCAAGCACAGTTTGGCCTGATGAAGAGG

Table B

Gene	Target Sequence Length	Amplicon Sequence
THBD.1	68	AGATCTGCGACGGACTGCGGGGCCACCTAATGACAGTGCGCTCCTCGGTGGCTGCCGATGTCATTTCC
THBS1.1	85	CATCCGCAAAGTGACTGAAGAGAACAAAGAGTTGGCCAATGAGCTGAGGCGGCCTCCCCTATGCTATCA CAACGGAGTTCAGTAC
TIMP1.1	76	TCCCTGCGGTCCCAGATAGCCTGAATCCTGCCCGGAGTGGAAGCTGAAGCCTGCACAGTGTCCACCCT GTTCCAC
TIMP2.1	69	TCACCCTCTGTGACTTCATCGTGCCCTGGGACACCCTGAGCACCACCCAGAAGAAGAGCCTGAACCACA
TIMP3.3	67	CTACCTGCCTTGCTTTGTGACTTCCAAGAACGAGTGTCTCTGGACCGACATGCTCTCCAATTTCCGT GCCGGAAGACCGTAATTGTGGCTGCACTGGATGGGACCTTCCAGAGGAAGCCATTTGGGGCCATCCT
TK1.2	84	GAACCTGGTGCCGCTG GGTTGGGCCACCTAGAAGTACTTGACCTGGGCCTTAATGAAATTGGGCAAGAAGTACAGGCCAGGAAT
TLR3.1	71	GG CCCTGAAAGAATGTTGTGGCTGCTCTTTTTCTGGTGACTGCCATTCATGCTGAACTCTGTCAACCAGGT
TMEM27.1	75	GCAGA GGATTCCACTGTTAGAGCCCTTACCGCCTGCTTATCCTACCCAATGACTACATTGGCTGTTGGTTATTTG
TMEM47.1	71	C
TMSB10.1	68	GAAATCGCCAGCTTCGATAAGGCCAAGCTGAAGAAAACGGAGACGCAGGAGAAGAACACCCTGCCGAC GGAGAAGGGTGACCGACTCAGCGCTGAGATCAATCGGCCCGACTATCTCGACTTTGCCGAGTCTGGGC
TNF.1	69	A
TNFAIP3.1	68	ATCGTCTTGGCTGAGAAAGGGAAAAGACACACAAGTCGCGTGGGTTGGAGAAGCCAGAGCCATTCCAC
TNFAIP6.1	67	AGGAGTGAAAGATGGGATGCCTATTGCTACAACCCACACGCAAAGGAGTGTGGTGGCGTCTTTACAG
TNFRSF10C.3	67	GGAGTTTGACCAGAGATGCAAGGGGTGAAGGAGCGCTTCCTACCGTTAGGGAACTCTGGGGACAGAG
TNFRSF10D.1	66	CCTCTCGCTTCTGGTGGTCTGTGAACTGAGTCCCTGGGATGCCTTTTAGGGCAGAGATTCCTGAGC
TNFRSF11B.1	67	TGGCGACCAAGACACCTTGAAGGGCCTAATGCACGCACTAAAGCACTCAAAGACGTACCACTTTCCC ACTGCCCTGAGCCCAAATGGGGGAGTGAGAGGCCATAGCTGTCTGGCATGGGCCTCTCCACCGTGCCT
TNFRSF1A.1	71	GAC
TNFSF12.1	68	TAGGCCAGGAGTTCCCAAATGTGAGGGGCGAGAAACAAGACAAGCTCCTCCCTTGAGAATTCCTGTG

Table B

Gene	Target Sequence Length	Amplicon Sequence
TNFSF13B.1	80	CCTACGCCATGGGACATCTAATTCAGAGGAAGAAGGTCCATGTCTTTGGGGATGAATTGAGTCTGGTGA CTTTGTTTCGA
TNFSF7.1	67	CCAACCTCACTGGGACACTTTTGCCTTCCCGAAACACTGATGAGACCTTCTTTGGAGTGCAGTGGGT
TNIP2.1	66	CATGTCAGAAAGGGCCGATCGGGAACGGGCTCAAAGTAGGATTCAAGAACTGGAGGAAAAGGTTCGC AATCCAAGGGGGAGAGTGATGACTTCCATATGGACTTTGACTCAGCTGTGGCTCCTCGGGCAAATCTG
TOP2A.4	72	TAC
TOP2B.2	66	TGTGGACATCTTCCCCTCAGACTTCCCTACTGAGCCACCTTCTCTGCCACGAACCGGTCTGGGCTAG CTATATGCAGCCAGAGATGTGACAGCCACCGTGGACAGCCTGCCACTCATCACAGCCTCCATTCTCAGT
TP.3	82	AAGAACTCGTGG CTTCACAGTGCTCCTGCAGTCTCTCTGTGTGGCTGTAACCTTACGTGTACTTTACCAACGAGCTGAAGCAG
TRAIL.1	73	ATG
TS.1	65	GCCTCGGTGTGCCTTTCAACATCGCCAGCTACGCCCTGCTCACGTACATGATTGCGCACATCACG
TSC1.1	66	TCACCAAATCTCAGCCCGCTTTCCTCATCGTTCAGCCGATGTCACCACCAGCCCTTATGCTGACAC
TSC2.1	69	CACAGTGGCCTCTTTCTCCTCCCTGTACCAGTCCAGCTGCCAAGGACAGCTGCACAGGAGCGTTTCCTG
TSPAN7.2	67	ATCACTGGGGTGATCCTGCTGGCTGTTGGAGTCTGGGGCAAACCTTACTCTGGGCACCTATATCTCCC CAGAAATCTCTGCAGGCAAGTTGCTCCAGAGCATATTGCAGGACAAGCCTGTAACGAATAGTTAAATTCA
TSPAN8.1	83	CGGCATCTGGATT
TUBB.1	73	CGAGGACGAGGCTTAAAAACTTCTCAGATCAATCGTGCATCCTTAGTGAACCTTCTGTTGTCCTCAAGCAT GGT
TUSC2.1	68	CACCAAGAACGGGCAGAAAGCGGGCCAAGCTGAGGCGAGTGCATAAGAATCTGATTCTCAGGGCATCG
tusc4.2	68	GGAGGAGCTAAATGCCTCAGGCCGGTGCACTCTGCCCATTGATGAGTCCAACACCATCCACTTGAAGG
TXLNA.1	68	GCCAGAACGGCTCAGTCTGGGGCCCTTCGTGATGTCTCTGAGGAGCTGAGCCGCCAACTGGAAGACAT

Table B

Gene	Target Sequence Length	Amplicon Sequence
UBB.1	522	GAGTCGACCCTGCACCTGGTCCCTGCGTCTGAGAGGTGGTATGCAGATCTTCGTGAAGACCCTGACCGG CAAGACCATCACCTGGAAGTGGAGCCAGTGACACCATCGAAAATGTGAAGGCCAAGATCCAGGATAA AGAAGGCATCCCTCCCGACCAGCAGAGGCTCATCTTTGCAGGCAAGCAGCTGGAAGATGGCCGCACTC TTTCTGACTACAACATCCAGAAGGAGTCGACCCTGCACCTGGTCCCTGCGTCTGAGAGGTGGTATGCAGA TCTTCGTGAAGACCCTGACCGGCAAGACCATCACTCTGGAAGTGGAGCCAGTGACACCATCGAAAATG TGAAGGCCAAGATCCAAGATAAAGAAGGCATCCCTCCCGACCAGCAGAGGCTCATCTTTGCAGGCAAGC AGCTGGAAGATGGCCGCACTCTTTCTGACTACAACATCCAGAAGGAGTCGACCCTGCACCTGGTCCCTGC GCCTGAGGGGTGGCTGTTAATTCTTCAGTCATGGCATTTCGC
UBE1C.1	76	TACCAG
UBE2C.1	67	TGTCTGGCGATAAAGGGATTTCTGCCTTCCCTGAATCAGACAACCTTTTCAAATGGGTAGGGACCAT
UBE2T.1	67	TGTTCTCAAATTGCCACCAAAAGGTGCTTGGAGACCATCCCTCAACATCGCAACTGTGTTGACCTCT GGCAACTGACAAACAGCCTTATAGCAAGCTCCCAGGTGTCTCTCTTCTGAAACCACTGAAAGGGGTAGAT
UGCG.1	73	CCT
UMOD.1	66	GCGTGGACCTGGATGAGTGCGCCATTCTGGAGCTCACAACCTGCTCCGCCAACAGCAGCTGCGTAA GTGGATGTGCCCTGAAGGACAAGCCAGGCGTCTACACGAGAGTCTCACACTTCTTACCCTGGATCCGCA
upa.3	70	G
USP34.1	70	AAGCTGTGATGGCCAAGCTTTGCCCTCCCAGGACCCTGAGGTTGCTTTATCTCTCAGTTGTGGCCATTC C
VCAM1.1	89	TGGCTTCAGGAGCTGAATACCCTCCCAGGCACACACAGGTGGGACACAAATAAGGGTTTTGGAACCACT ATTTTCTCATCACGACAGCA
VCAN.1	69	CCTGCTACACAGCCAACAAGACCACCCACTGTGGAAGACAAAGAGGCCTTTGGACCTCAGGCGCTTTCT
VDR.2	67	GCCCTGGATTTTCAAGAAAGAGCCAAGTCTGGATCTGGGACCCTTTCCTTCCCTGGCTTGTAAC CTGCTGTCTTGGGTGCATTGGAGCCTTGCTTGCTCTACCTCCACCATGCCAAGTGGTCCCAGGCT
VEGF.1	71	GC
VEGFB.1	71	TGACGATGGCCTGGAGTGTGTGCCCACTGGGCAGCACCAAGTCCGGATGCAGATCCTCATGATCCGGT ACC
VHL.1	67	CGGTTGGTGACTTGTCTGCCTCCTGCTTTGGGAAGACTGAGGCATCCGTGAGGCAGGGACAAGTCTT TGCCCTTAAAGGAACCAATGAGTCCCTGGAACGCCAGATGCGTGAAATGGAAGAGAAGTCTTGCCGTTGA
VIM.3	72	AGC
VTCN1.1	70	ACAGTGGTCTGGGCATCCCAAGTTGACCAGGGAGCCAACCTTCTCGGAAGTCTCCAATACCAGCTTTGAG C
VTN.1	67	AGTCAATCTTCGCACACGGCGAGTGGACACTGTGGACCCTCCCTACCCACGCTCCATCGCTCAGTAC

Table B

Gene	Target Sequence Length	Amplicon Sequence
VWF.1	66	TGAAGCACAGTGCCCTCTCCGTCGAGCTGCACAGTGACATGGAGGTGACGGTGAATGGGAGACTGG
WIF.1	67	AACAAGCTGAGTGCCCAGGCGGGTGCCGAAATGGAGGCTTTTGTAAATGAAAGACGCATCTGCGAGTG AGAGGCATCCATGAACTTCACACTTGCGGGCTGCATCAGCACACGCTCCTATCAACCCAAGTACTGTGG
WISP1.1	75	AGTTTG
WT1.1	66	TGTACGGTCGGCATCTGAGACCAGTGAGAAACGCCCCTTCATGTGTGCTTACCCAGGCTGCAATAA ATCGCAGCTGGTGGGTGTACACACTGCTGTTTACCTTGGCGAGGCCTTTCACCAAGTCCATGCAACAGG
WWOX.5	74	GAGCT
XDH.1	66	TGGTGGCAGACATCCCTTCCTGGCCAGATACAAGGTTGGCTTCATGAAGACTGGGACAGTTGTGGC GCAGTTGGAAGACACAGGAAAGTATCCCCAAATTGCAGATTTATCAACGGCTTTTATCTTGAAAATAGTG
XIAP.1	77	CCACGCA CACCTGCACTGAACATACCCCAAGAGCCCCTGCTGGCCCATTGCCTAGAAACCTTTGCATTCATCCTCC
XPNPEP2.2	72	TT AGACTGTGGAGTTTGATGTTGTTGAAGGAGAAAAGGGTGCGGAGGCAGCAAATGTTACAGGTCCTGGTG
YB-1.2	76	GTGTTCC
ZHX2.1	67	GAGTACGACCAGTTAGCGGCCAAGACTGGCCTGGTCCGAACTGAGATTGTGCGTTGGTTCAAGGAGA

CLAIMS

What is claimed is:

1. A method for determining a likely clinical outcome for a patient with cancer comprising:
 - (a) measuring, in a biological sample obtained from the patient, an expression level of at least one gene, or its expression product, from Tables 3a, 3b, 8a or 8b;
 - (b) normalizing the expression level to obtain a normalized expression level; and
 - (c) calculating a score estimating the likelihood that the patient will have a negative clinical outcome based on the normalized expression level,wherein expression of genes listed in Tables 3a and 8a are negatively correlated with a risk of a negative clinical outcome, and
wherein expression of genes listed in Tables 3b and 8b are positively correlated with a risk of a negative clinical outcome.
2. The method of claim 1, further comprising generating a report based on the score.
3. The method of claim 1, wherein the negative clinical outcome is a recurrence of cancer.
4. The method of claim 3, where the biological sample is obtained from a tumor.
5. The method of claim 4, wherein the tumor is a renal cell carcinoma.
6. The method of claim 1, wherein expression levels from at least two genes are measured.
7. The method of claim 6, wherein the at least two genes are from one or more gene subsets.
8. The method of claim 7, wherein the one or more gene subsets is angiogenesis, immune response, transport, cell adhesion/extracellular matrix, cell cycle, or apoptosis genes.

9. A method for determining stage for a tumor obtained from a cancer patient, comprising:
 - (a) measuring, in a biological sample obtained from the tumor, an expression level of at least one proxy gene, or its gene product, from Tables 4a or 4b;
 - (b) normalizing the expression level to obtain a normalized expression level; and
 - (c) determining the stage of the tumor based on the normalized expression level, wherein expression of proxy genes listed in Table 4a is positively correlated with increased stage of the tumor, and wherein expression of proxy genes listed in Table 4b is negatively correlated with increased stage of the tumor.
10. The method of claim 9, wherein the tumor is renal cell carcinoma.
11. A method for determining grade of a tumor obtained from a cancer patient, comprising:
 - (a) measuring, in a biological sample obtained from the tumor, an expression level of at least one proxy gene, or its gene product, from Tables 5a and 5b;
 - (b) normalizing the expression level to obtain a normalized expression level; and
 - (c) determining grade of the tumor based on the normalized expression level, wherein expression of proxy genes listed in Table 5a is positively correlated with increased grade of the tumor, and wherein expression of proxy genes listed in Table 5b is negatively correlated with increased grade of the tumor.
12. The method of claim 11, wherein the tumor is renal cell carcinoma.

13. A method for determining presence of necrosis in a tissue sample obtained from a patient with cancer, comprising:

(a) measuring, in the tissue sample, an expression level of at least one proxy gene, or its gene product, from Tables 6a and 6b;

(b) normalizing the expression level to obtain a normalized expression level; and

(c) determining the presence of necrosis of the tissue based on the normalized expression level,

wherein expression of proxy genes listed in Table 6a is positively correlated with the presence of necrosis, and

wherein expression of proxy genes listed in Table 6b is negatively correlated with the presence of necrosis.

14. The method of claim 13, wherein the cancer is renal cell carcinoma.

15. A method for determining nodal status of a patient with cancer, comprising:

(a) measuring, in a biological sample obtained from the cancer patient, an expression level of at least one proxy gene, or its gene product, from Tables 7a and 7b;

(b) normalizing the expression level to obtain a normalized expression level; and

(c) determining the nodal status of the cancer patient based on the normalized expression level,

wherein expression of proxy genes listed in Table 7a is positively correlated with nodal invasion, and

wherein expression of proxy genes listed in Table 7b is negatively correlated with nodal invasion.

16. The method of claim 15, wherein the cancer is renal cell carcinoma.