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(54) **LARGE-SCALE EPIGENOMIC REPROGRAMMING LINKS ANABOLIC GLUCOSE METABOLISM TO DISTANT METASTASIS DURING THE EVOLUTION OF PANCREATIC CANCER PROGRESSION**

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Related U.S. Application Data

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(60) Provisional application No. 62/405,155, filed on Oct. 6, 2016.

Publication Classification

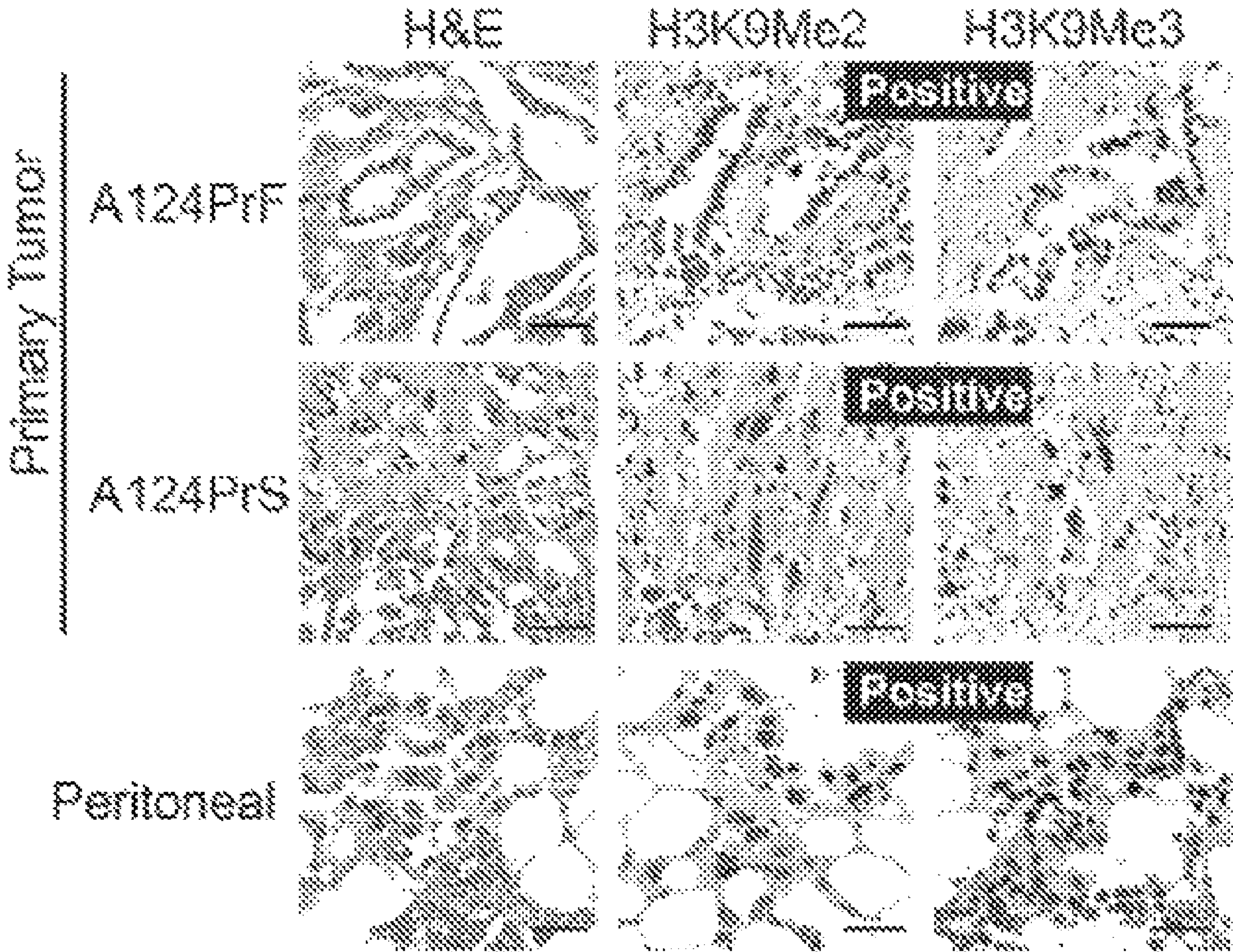
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C12Q 1/6874 (2006.01)

(52) **U.S. Cl.**
CPC **C12Q 1/6811** (2013.01); **C12Q 1/6874** (2013.01); **C12Q 2600/158** (2013.01); **C12Q 2600/112** (2013.01); **C12Q 1/6806** (2013.01)

(57) **ABSTRACT**

The present invention relates to a method of identifying epigenetic reprogramming. Identifying epigenetic reprogramming comprises detecting large organized heterochromatin lysine (K)-9 modified domains (LOCKS) and large DNA hypomethylated blocks in a sample containing DNA from a subject having cancer, for example, PDAC.

Specification includes a Sequence Listing.



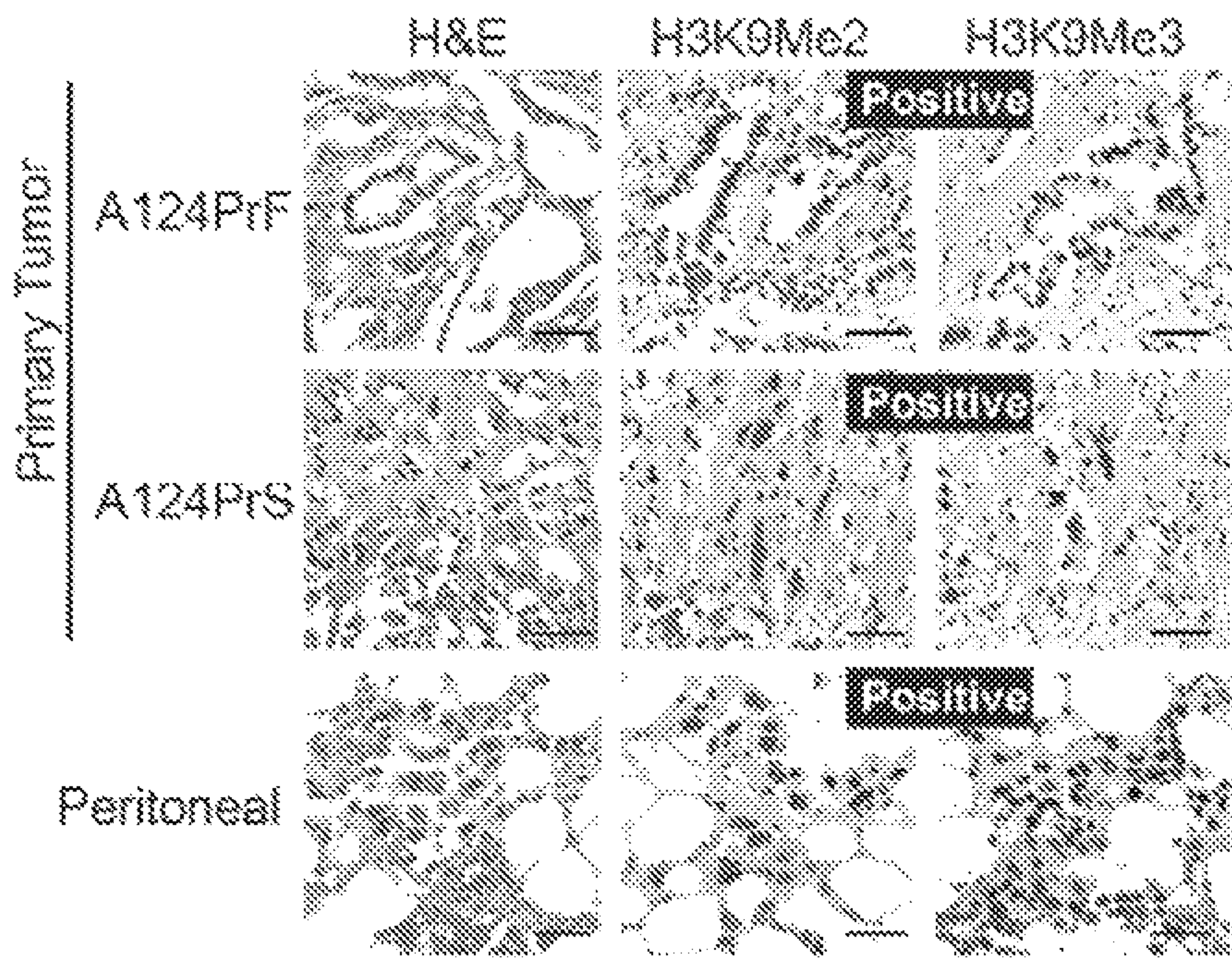


FIGURE 1A

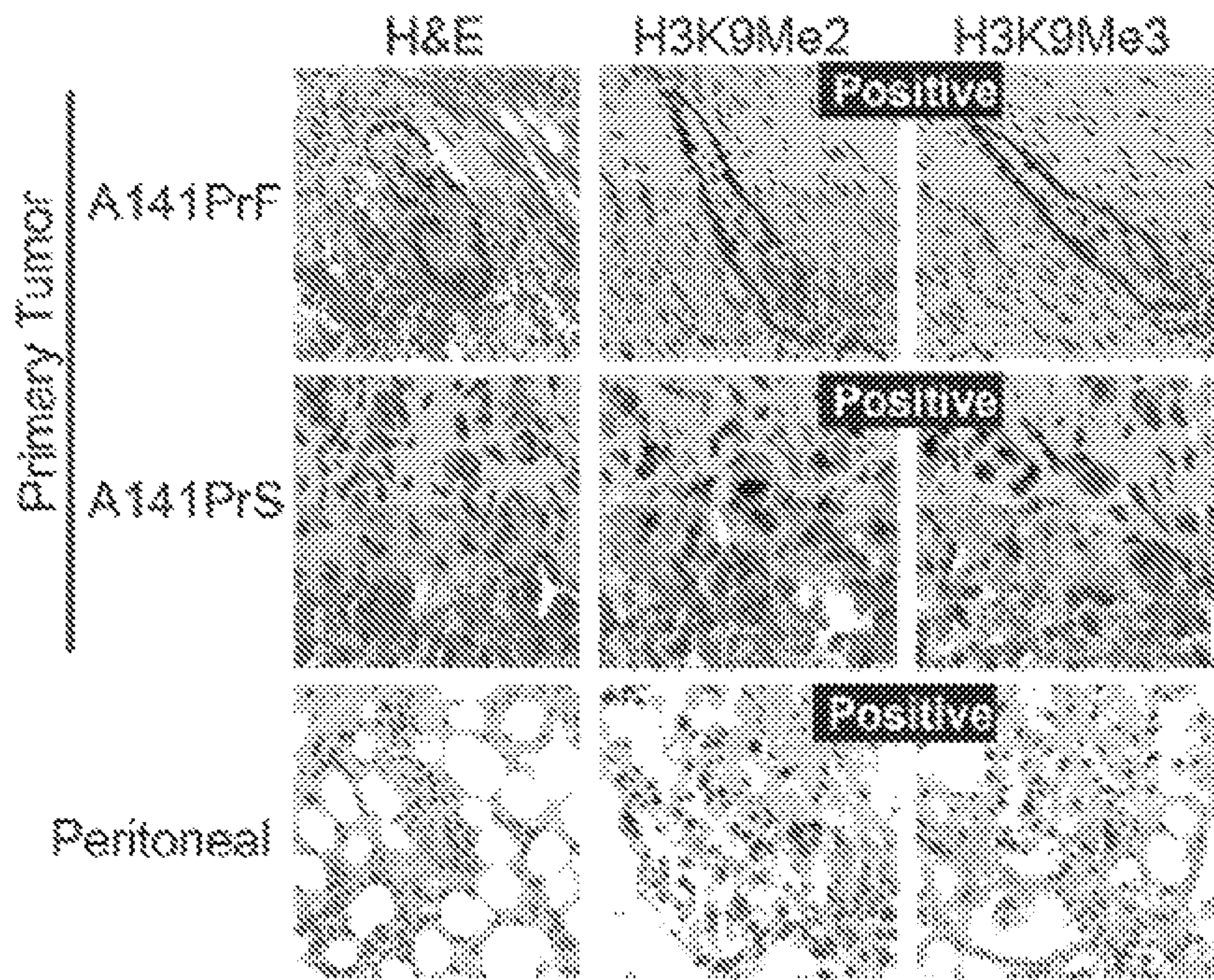


FIGURE 1B

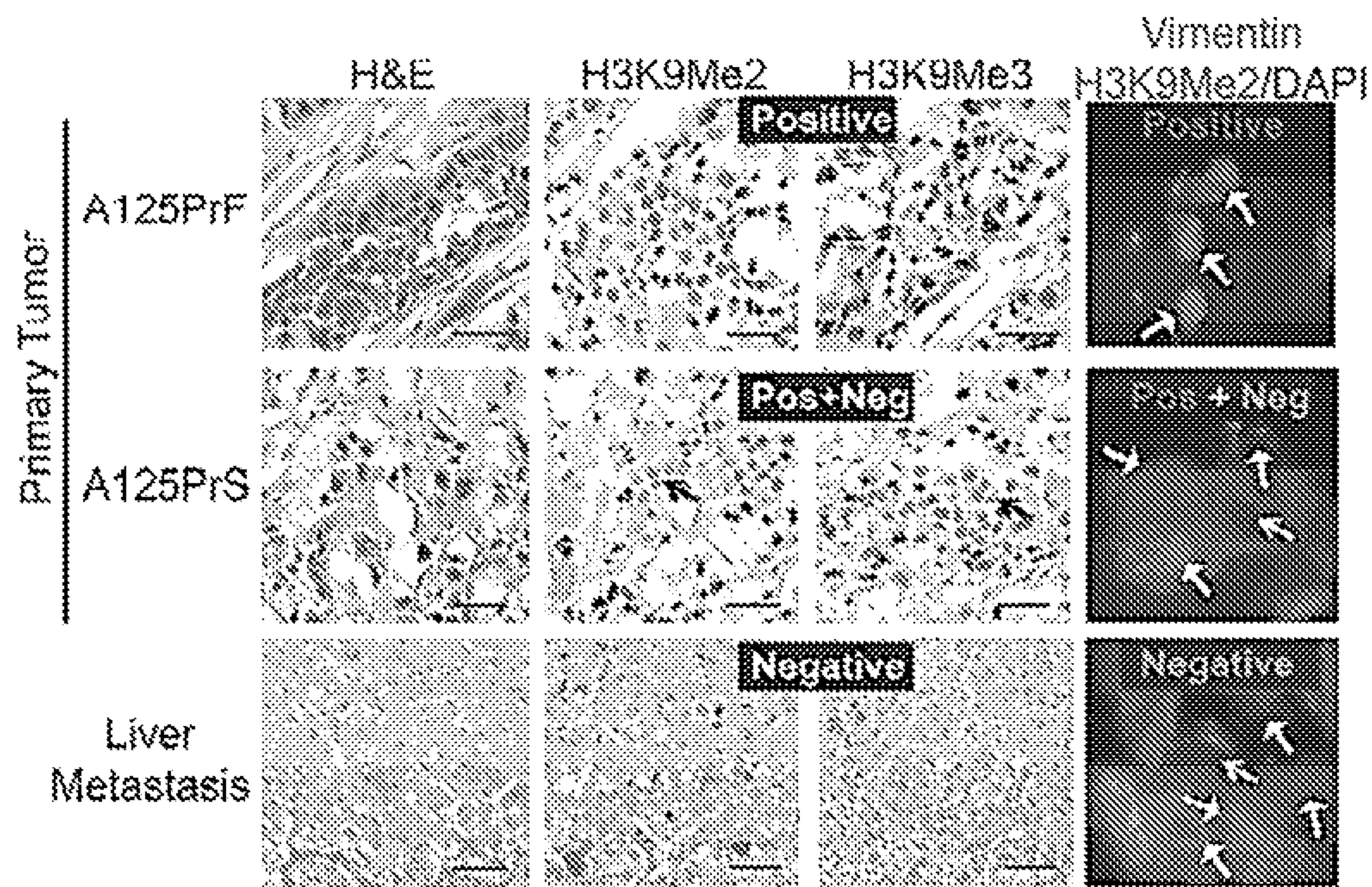


FIGURE 1C

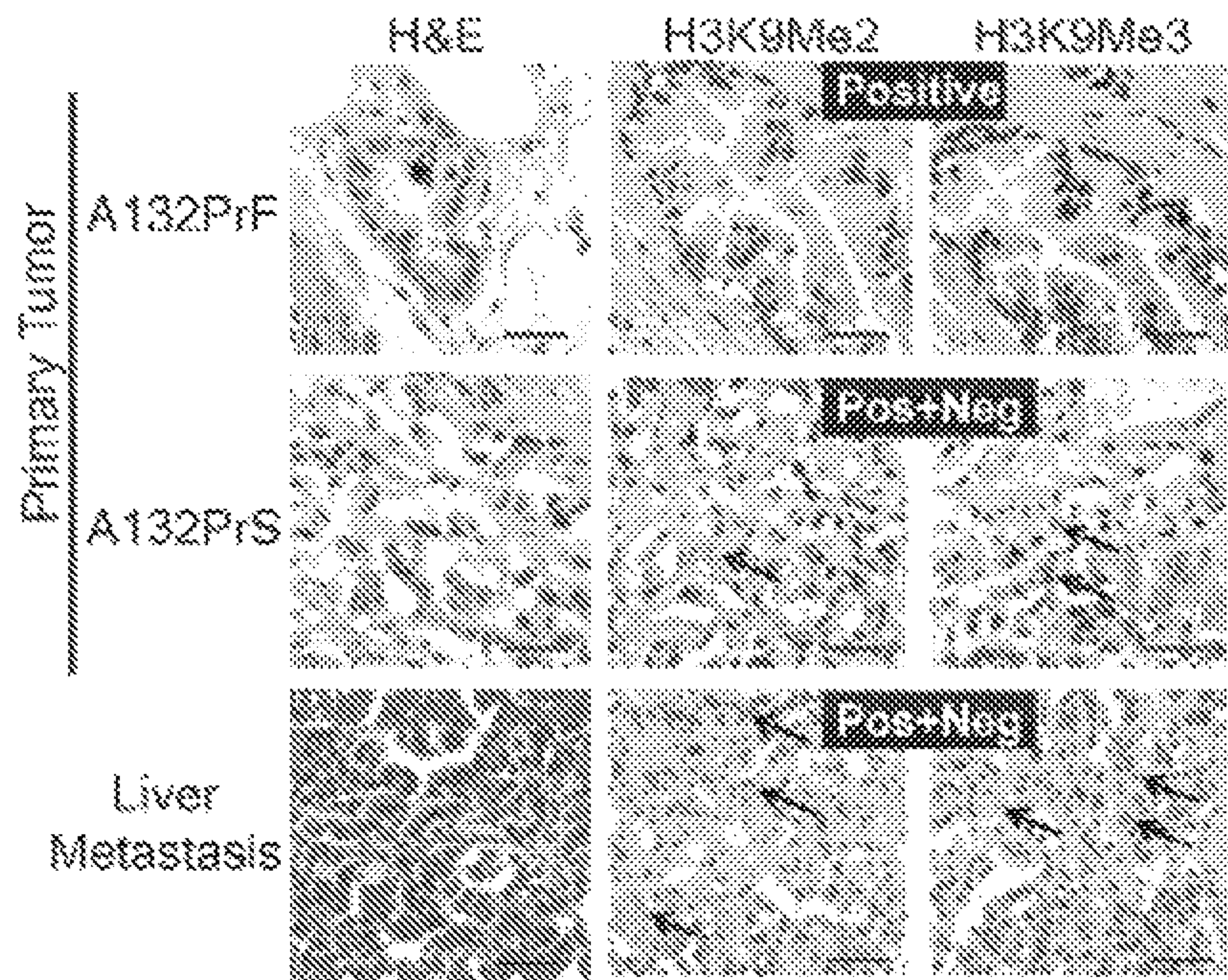


FIGURE 1D

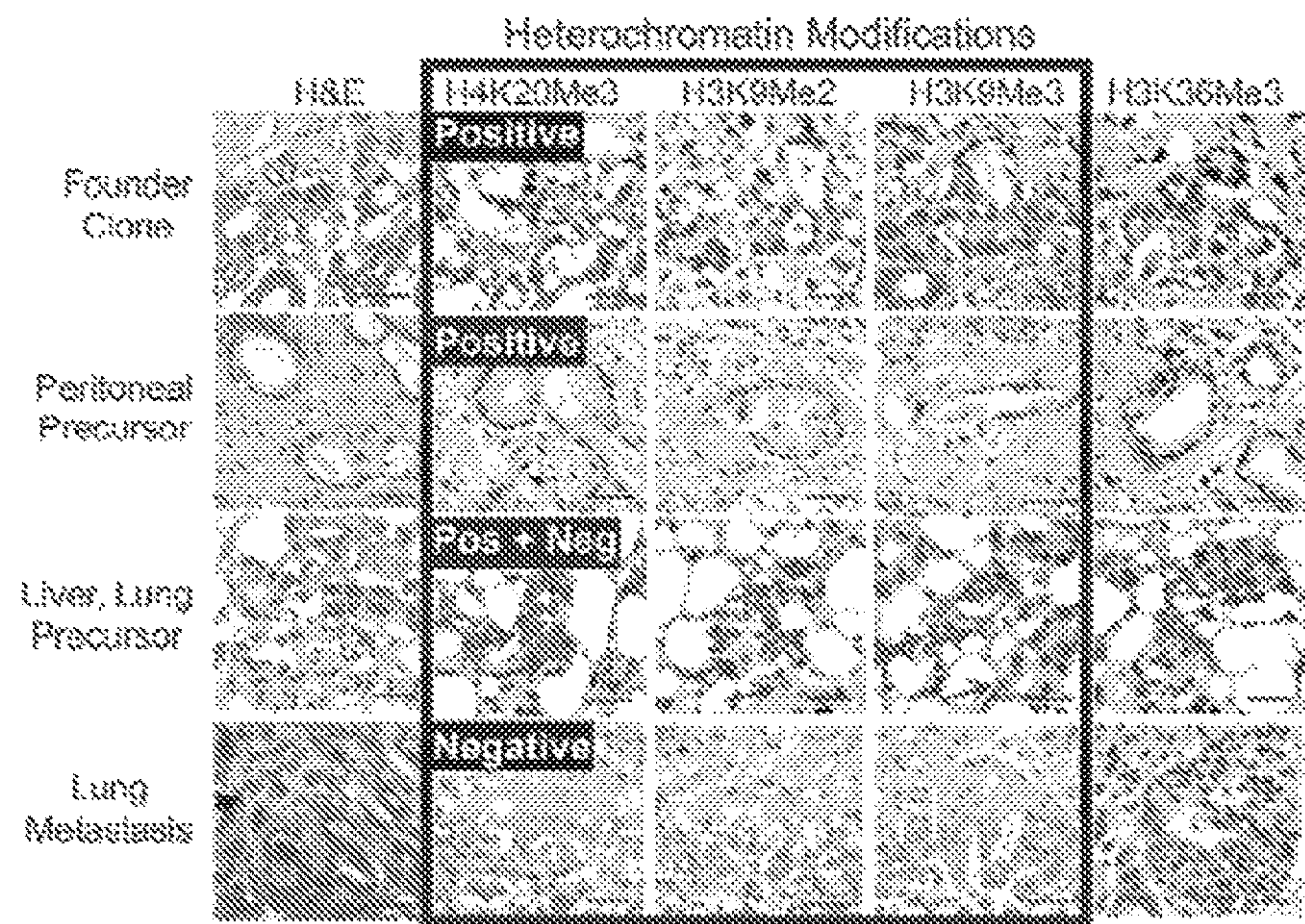


FIGURE 1E

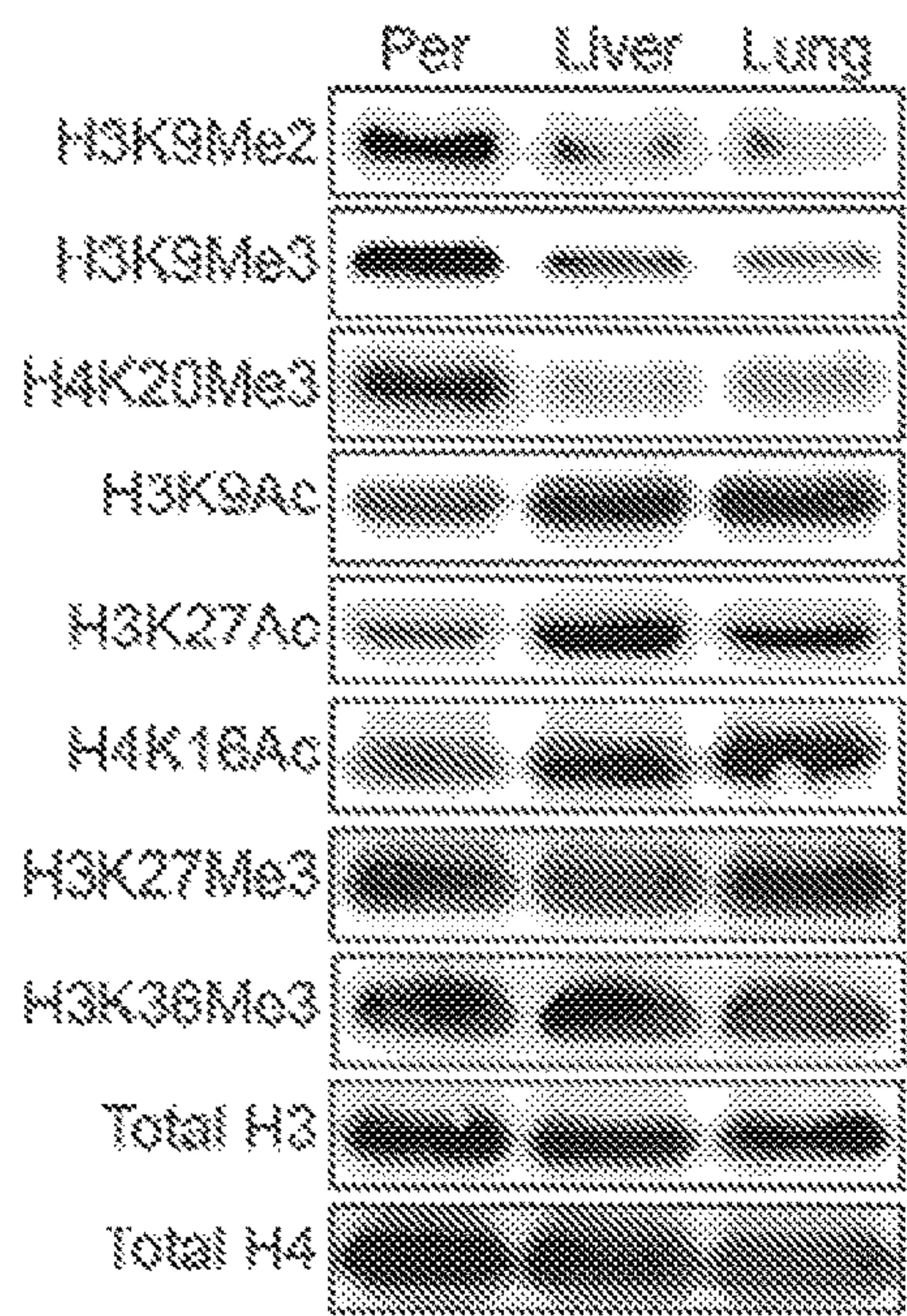


FIGURE 1F

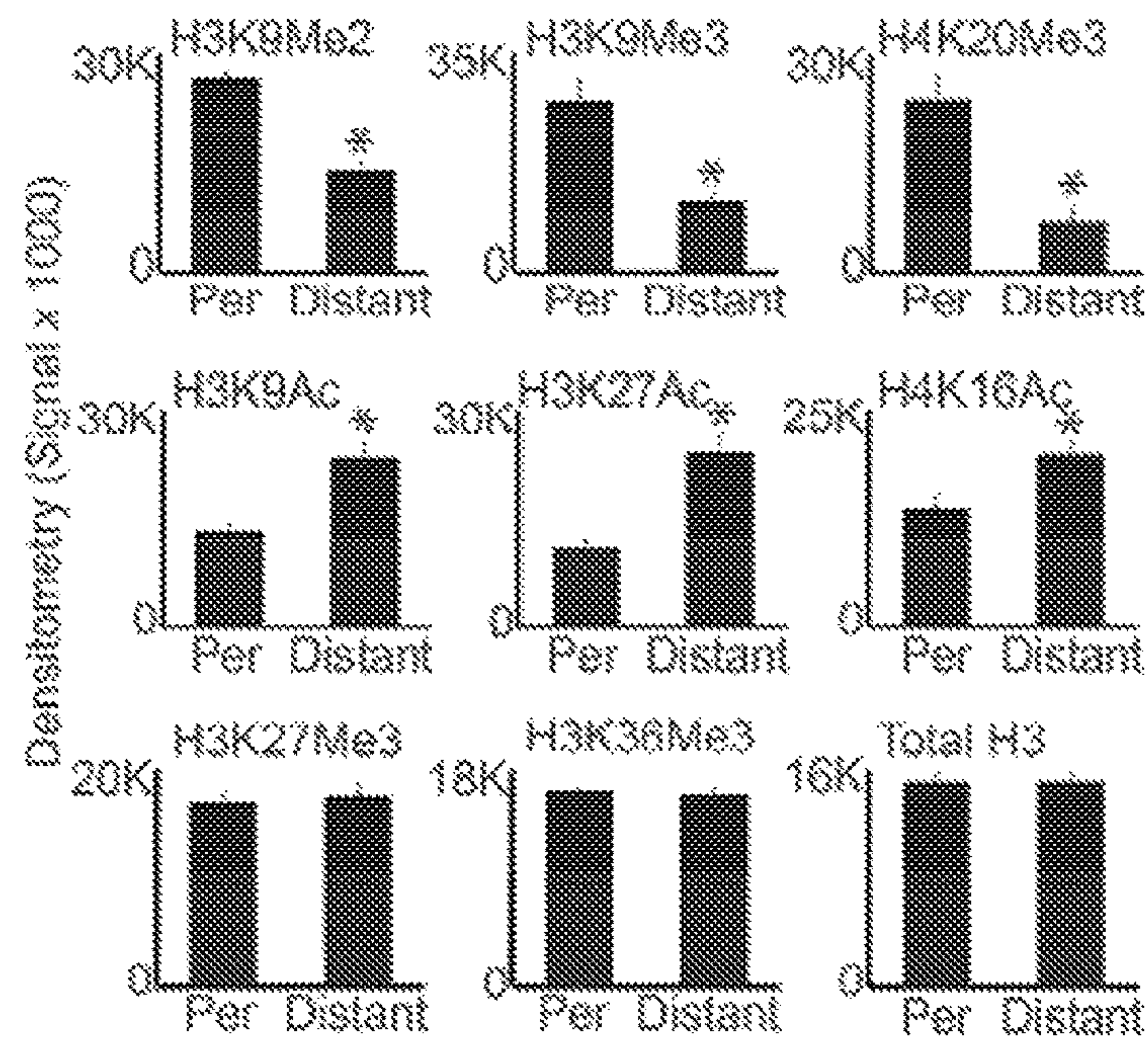


FIGURE 1G

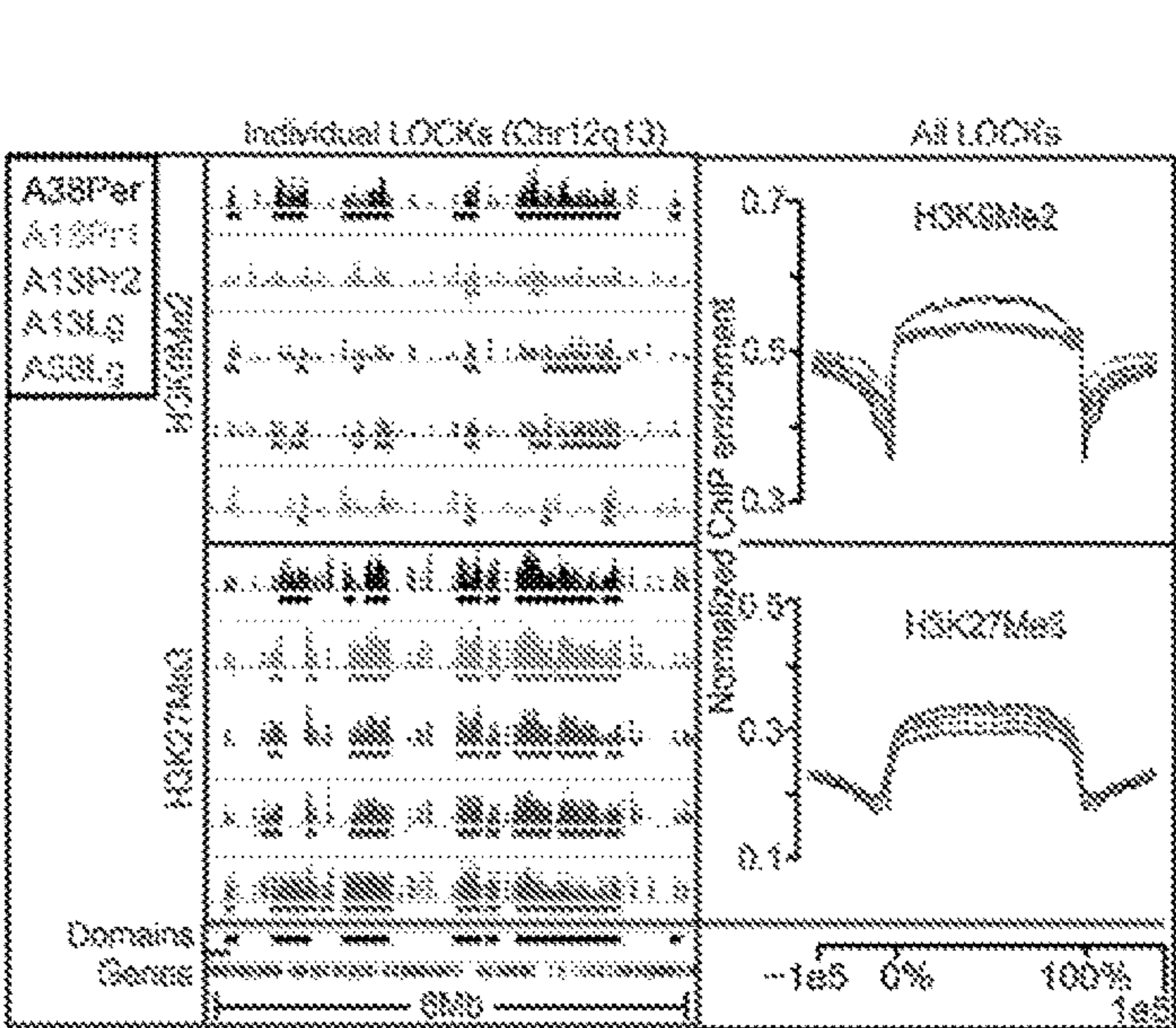


FIGURE 2A

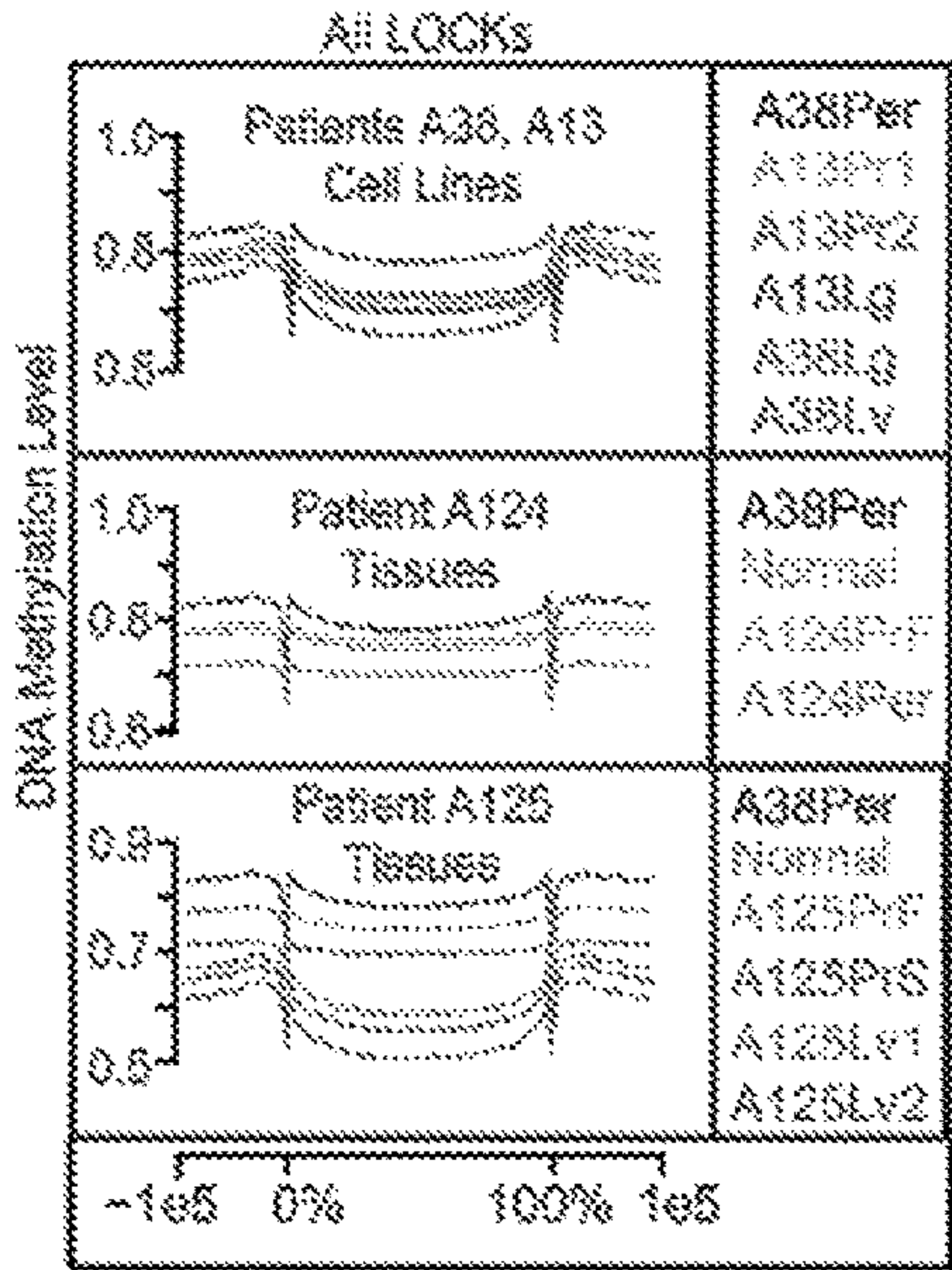


FIGURE 2B

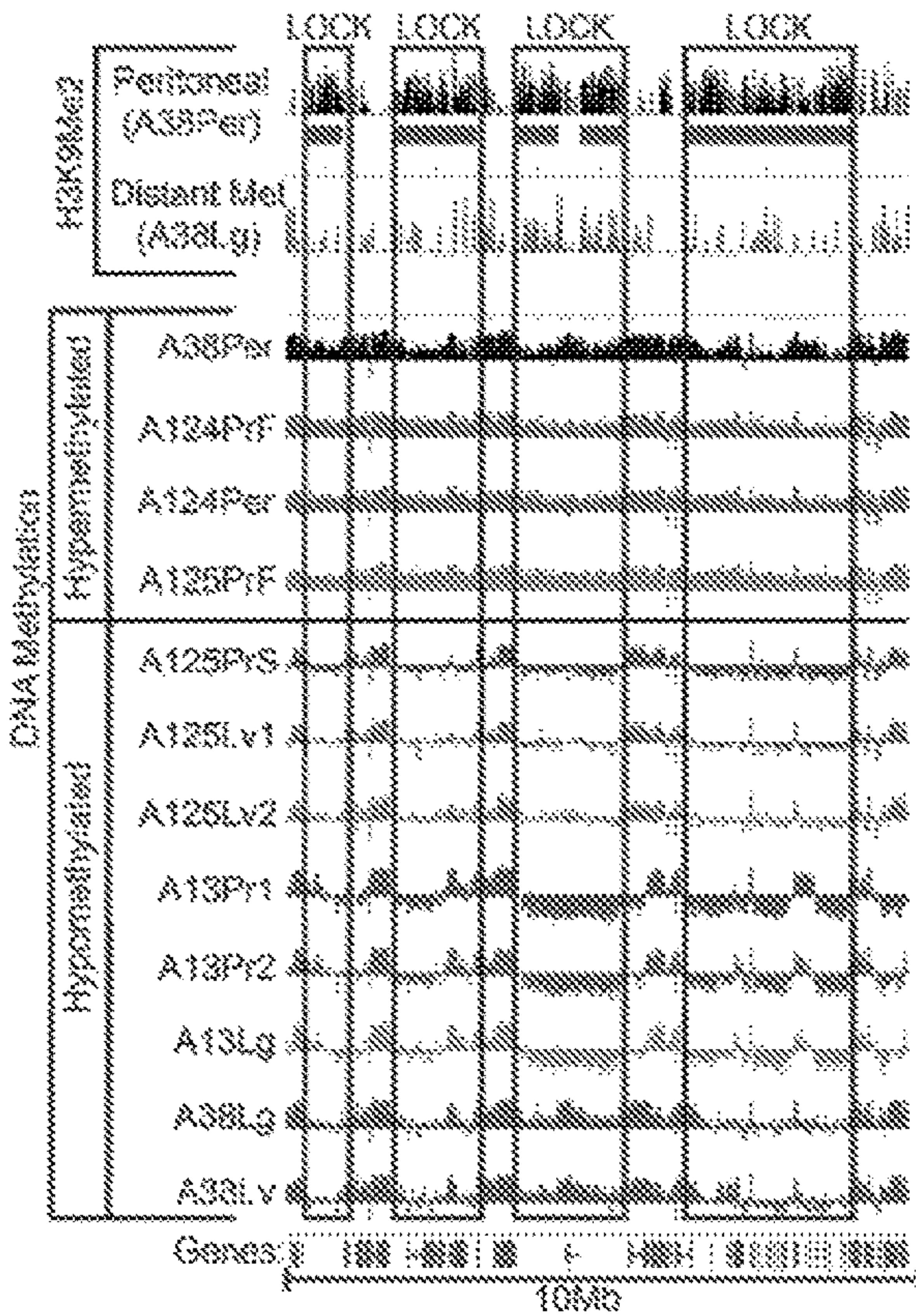


FIGURE 2C

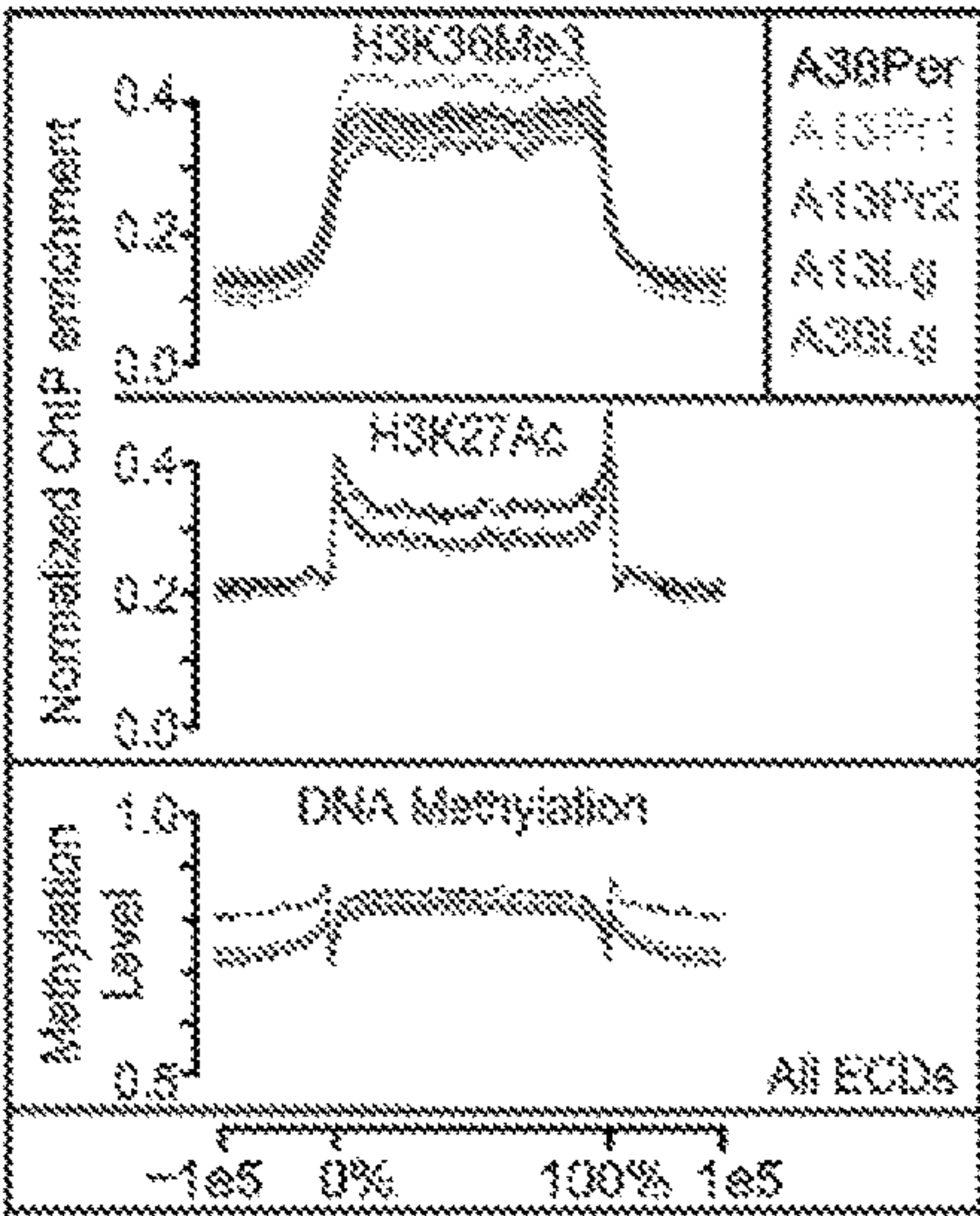


FIGURE 2D

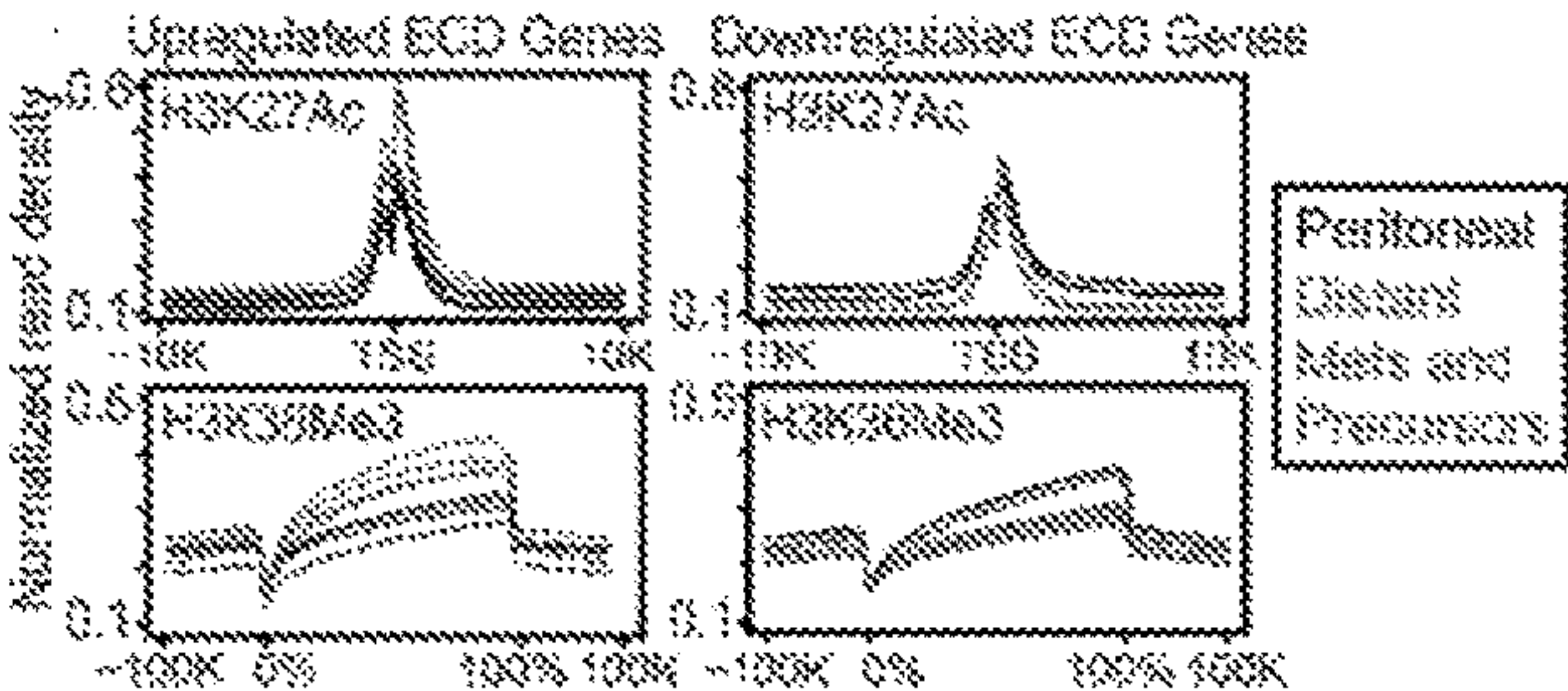


FIGURE 2E

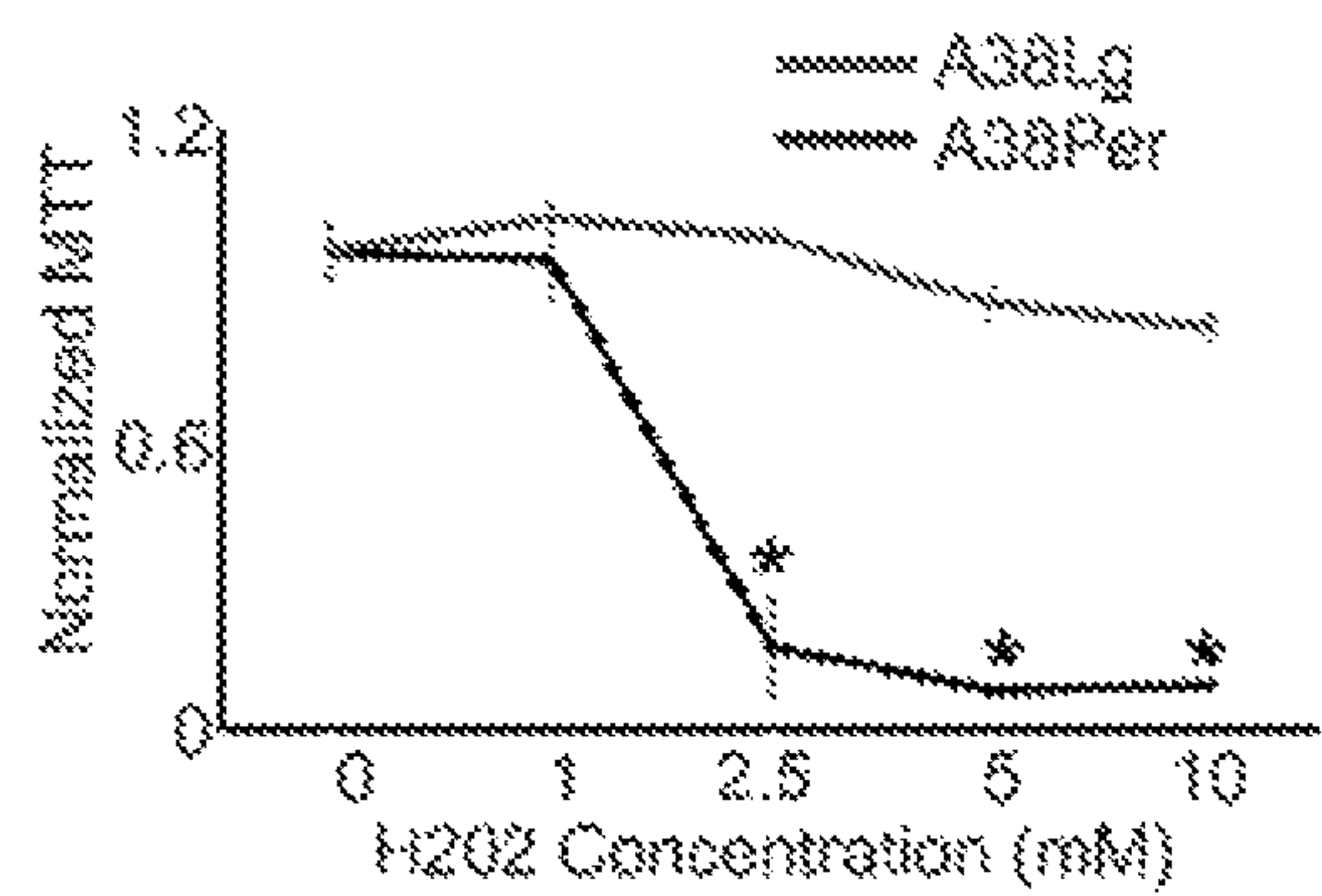


FIGURE 3A

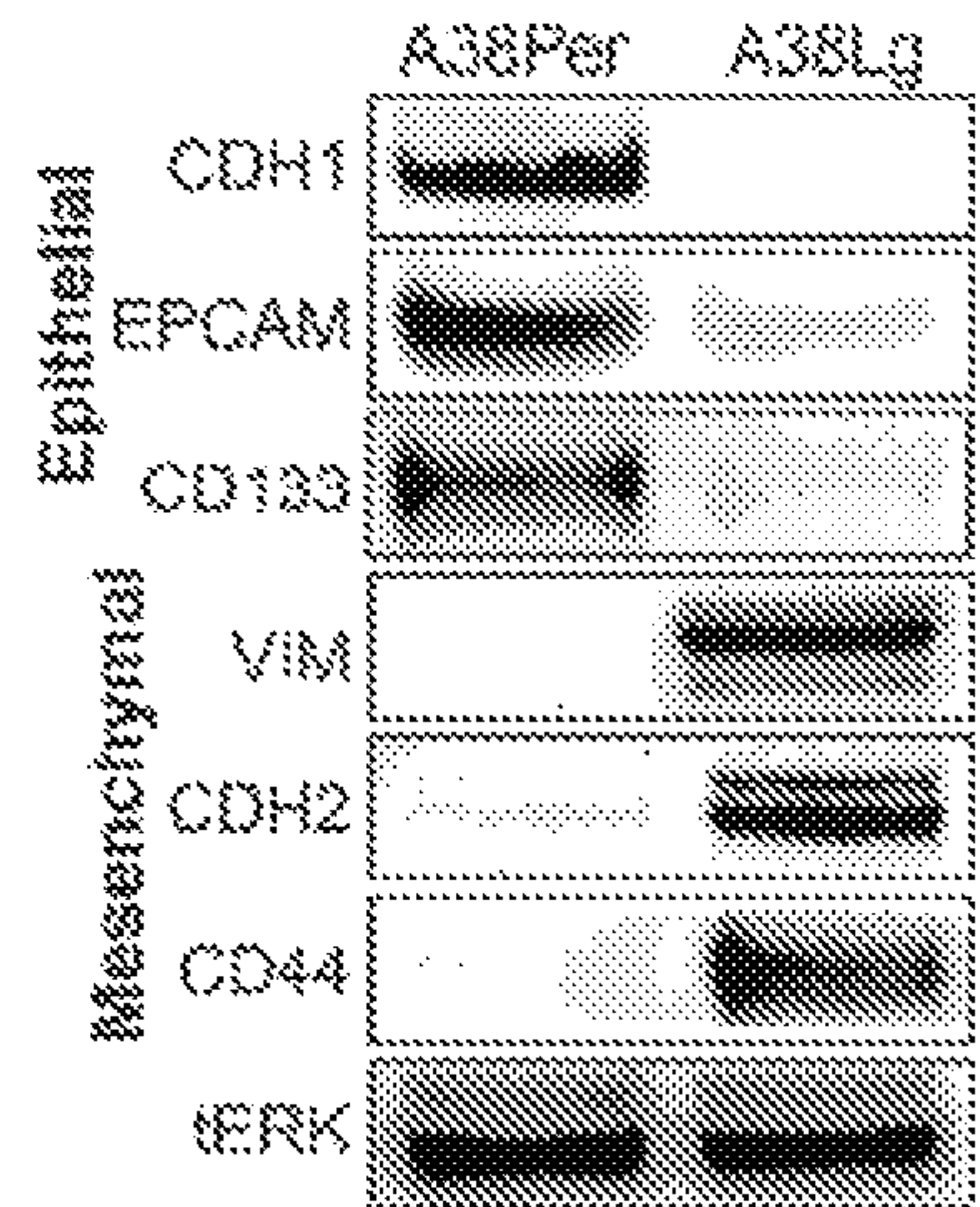


FIGURE 3B

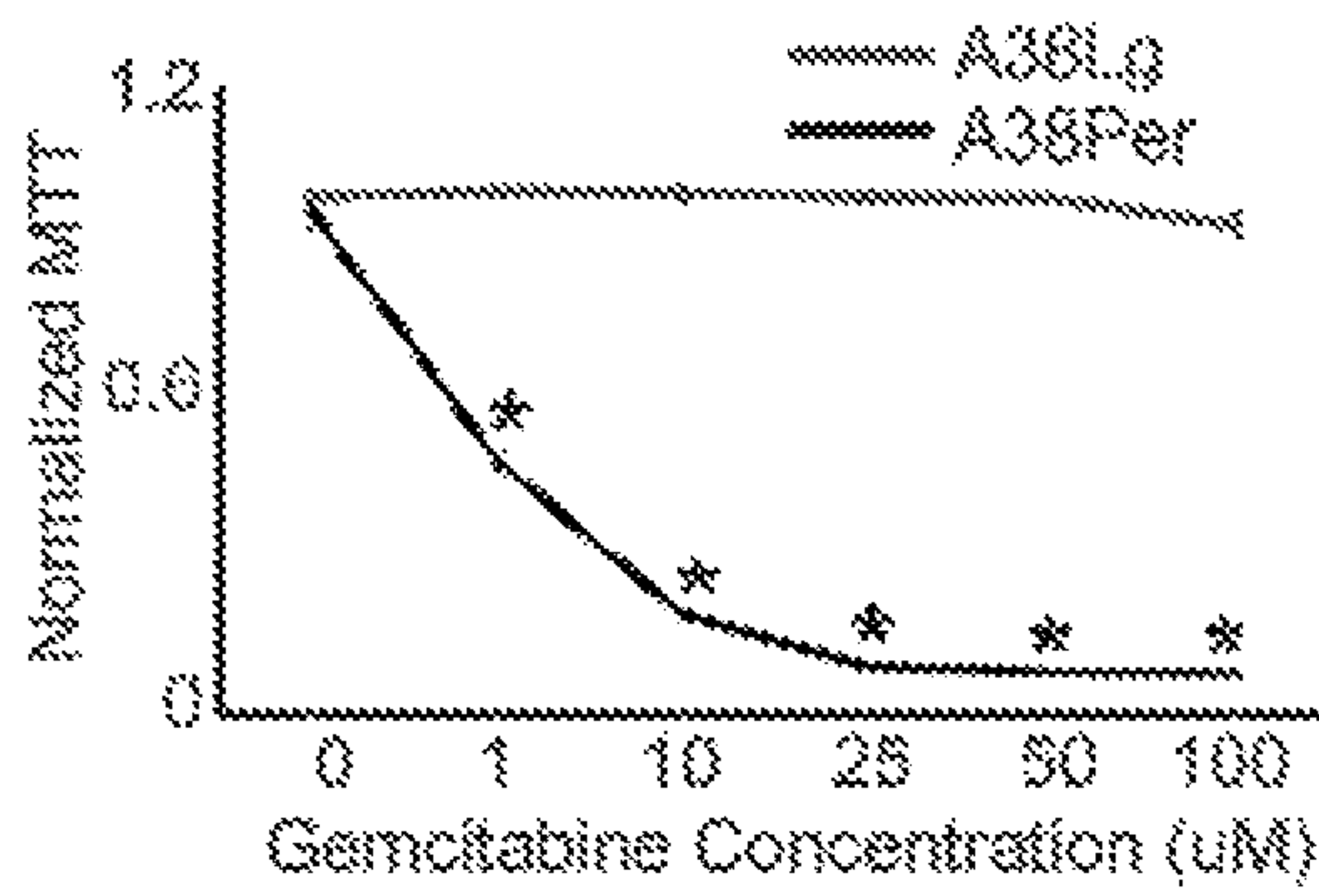


FIGURE 3C

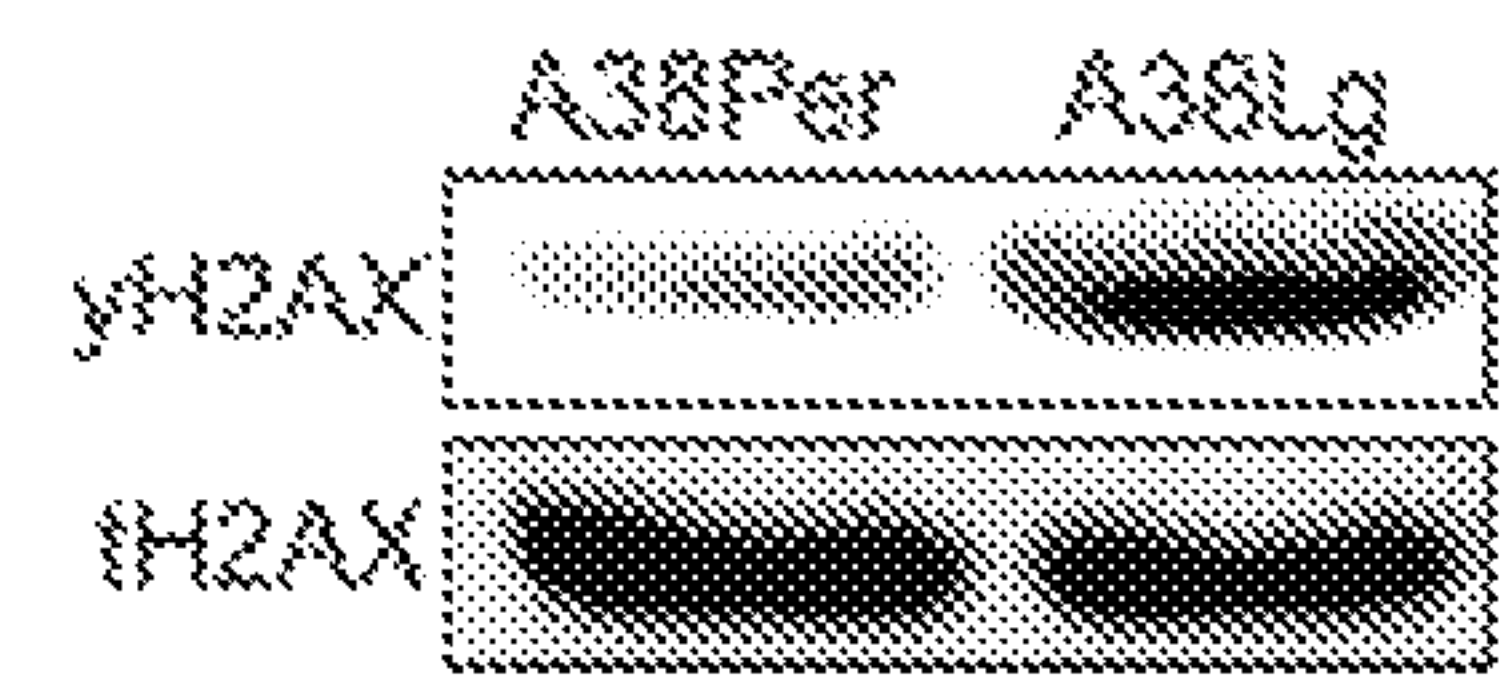


FIGURE 3D

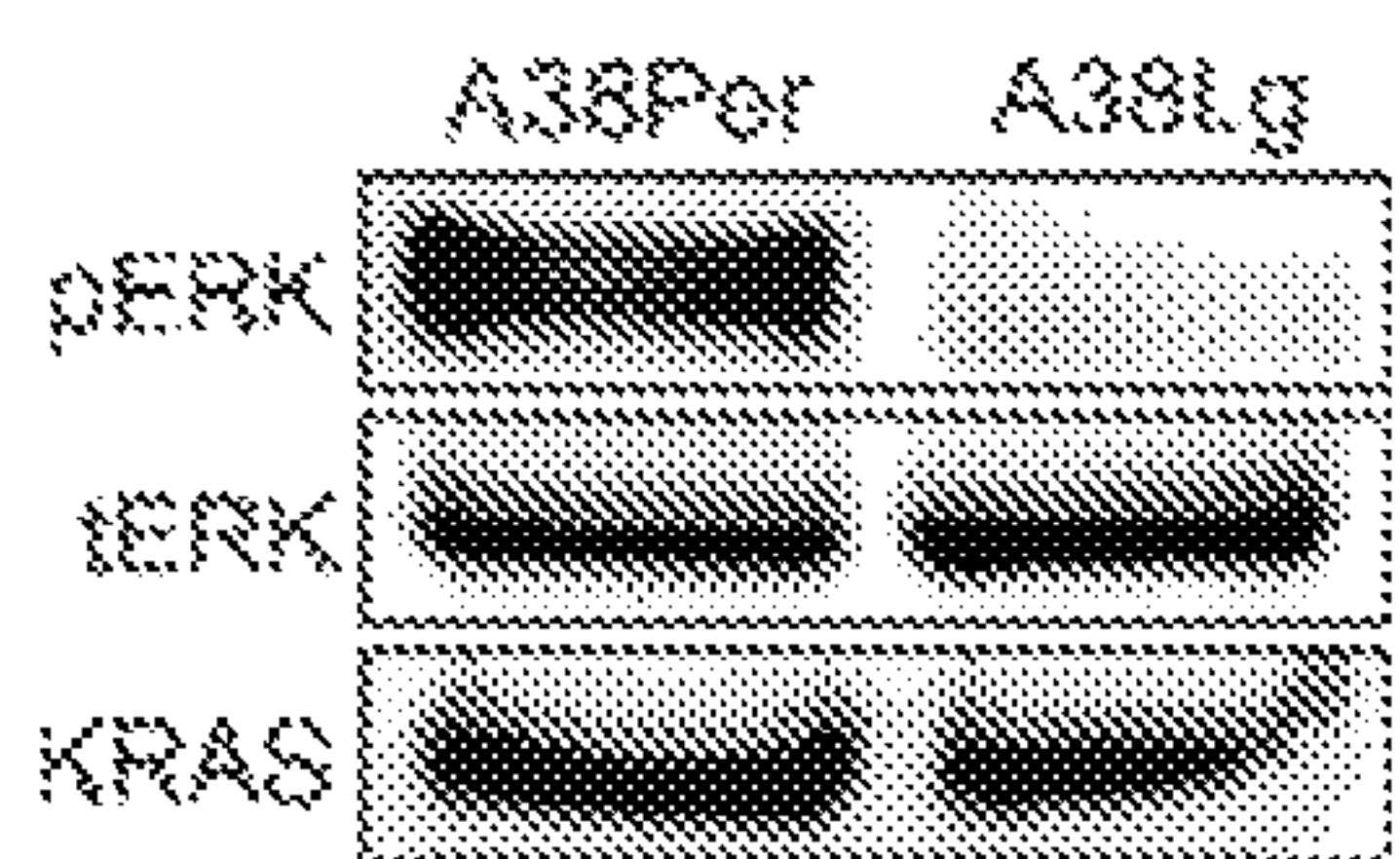


FIGURE 3E

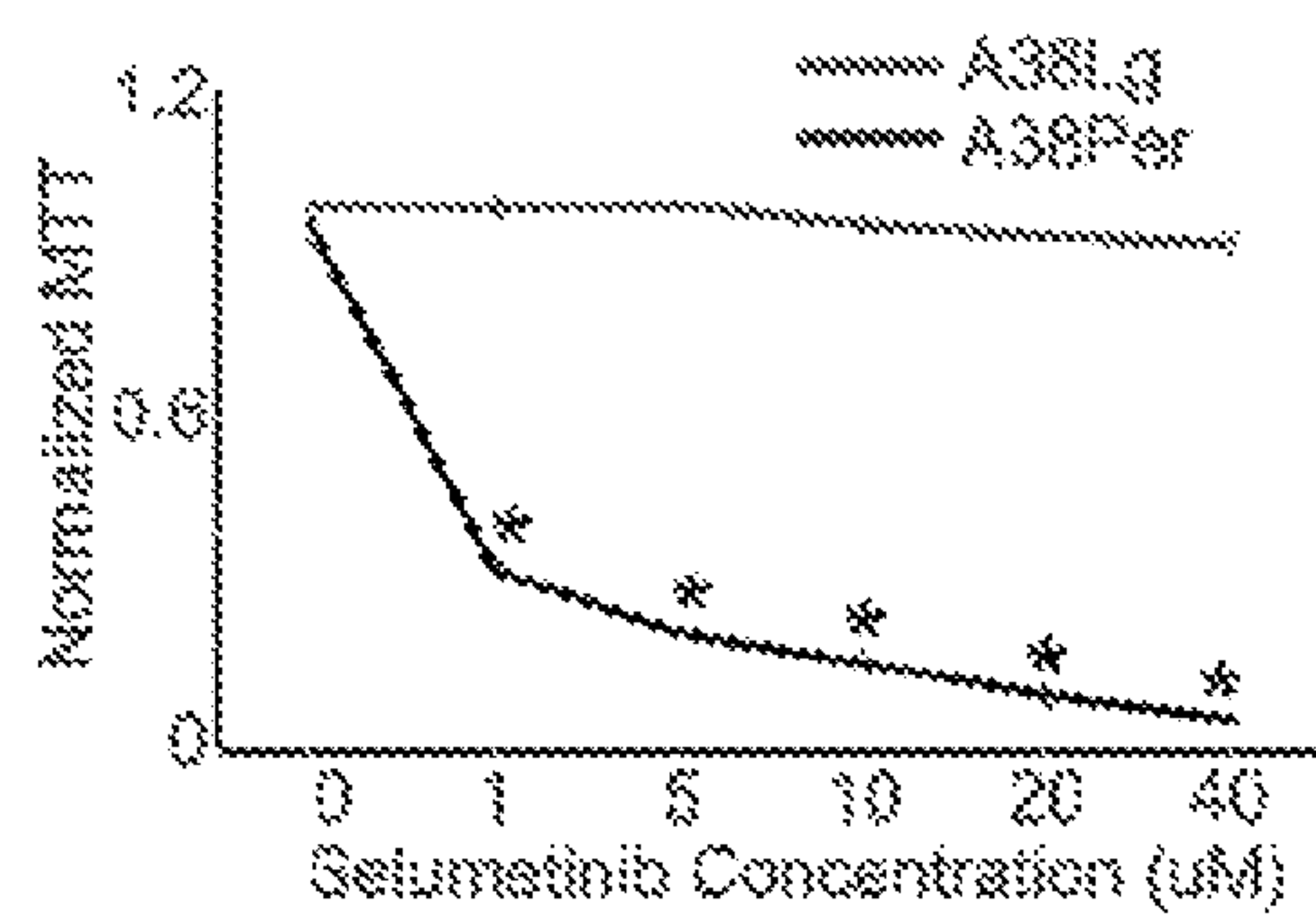


FIGURE 3F

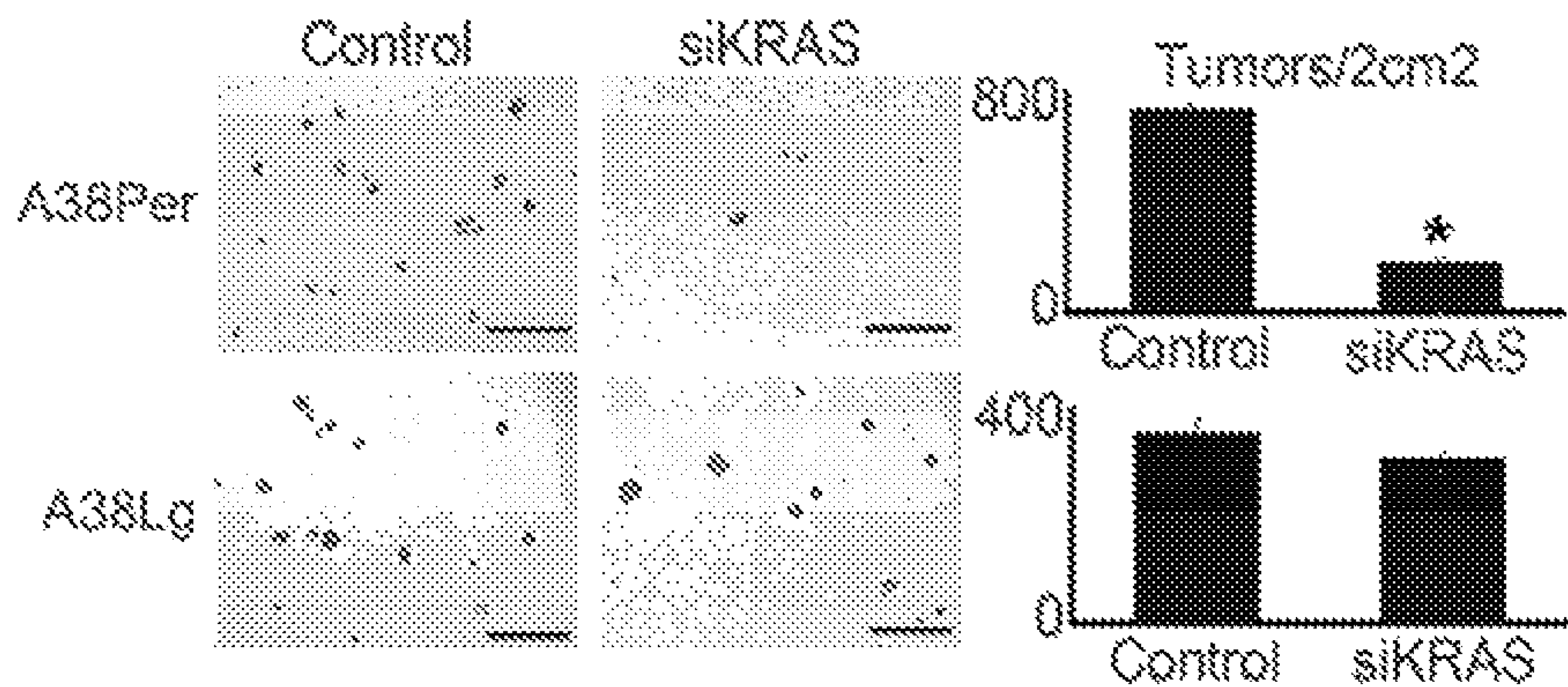
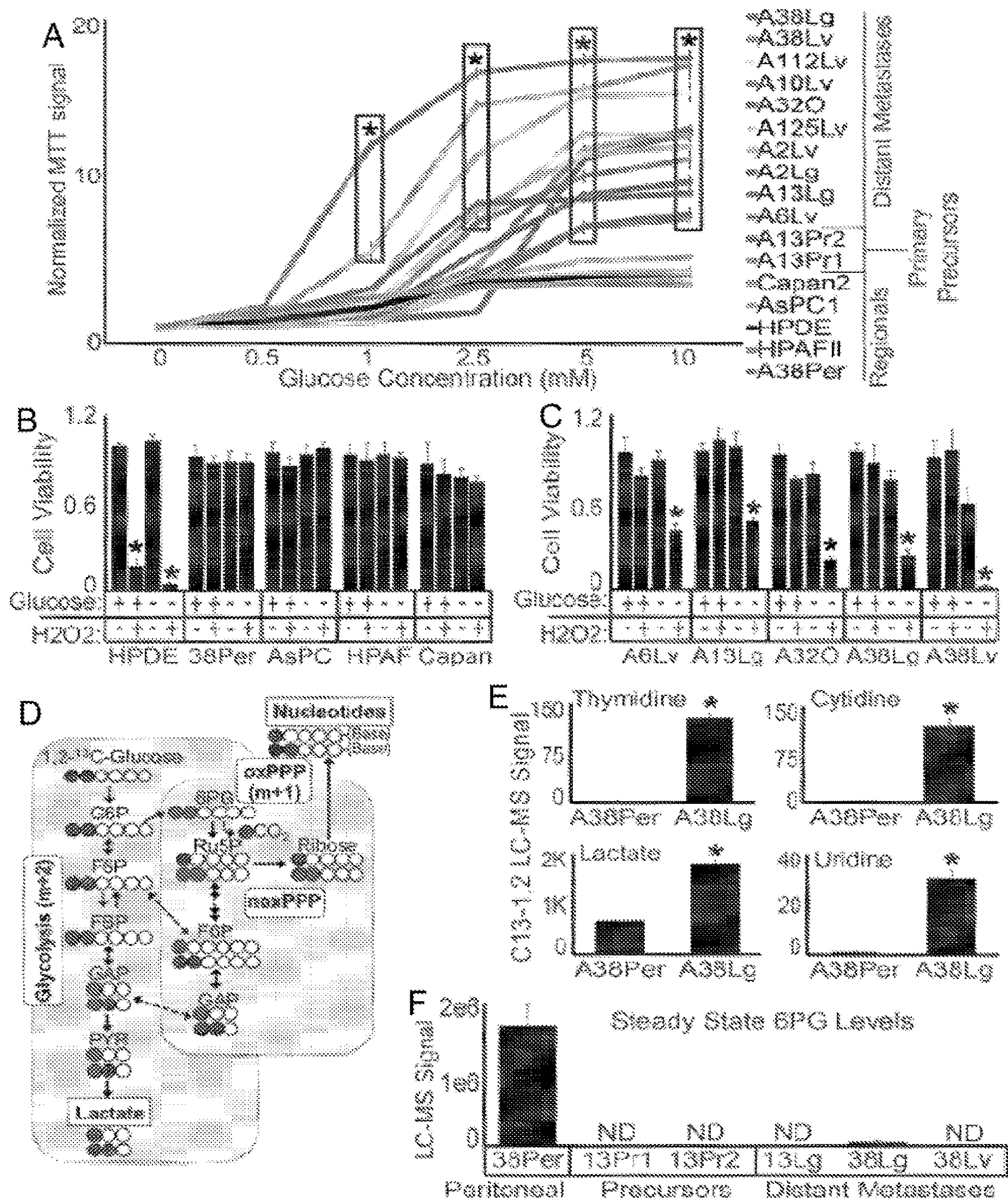
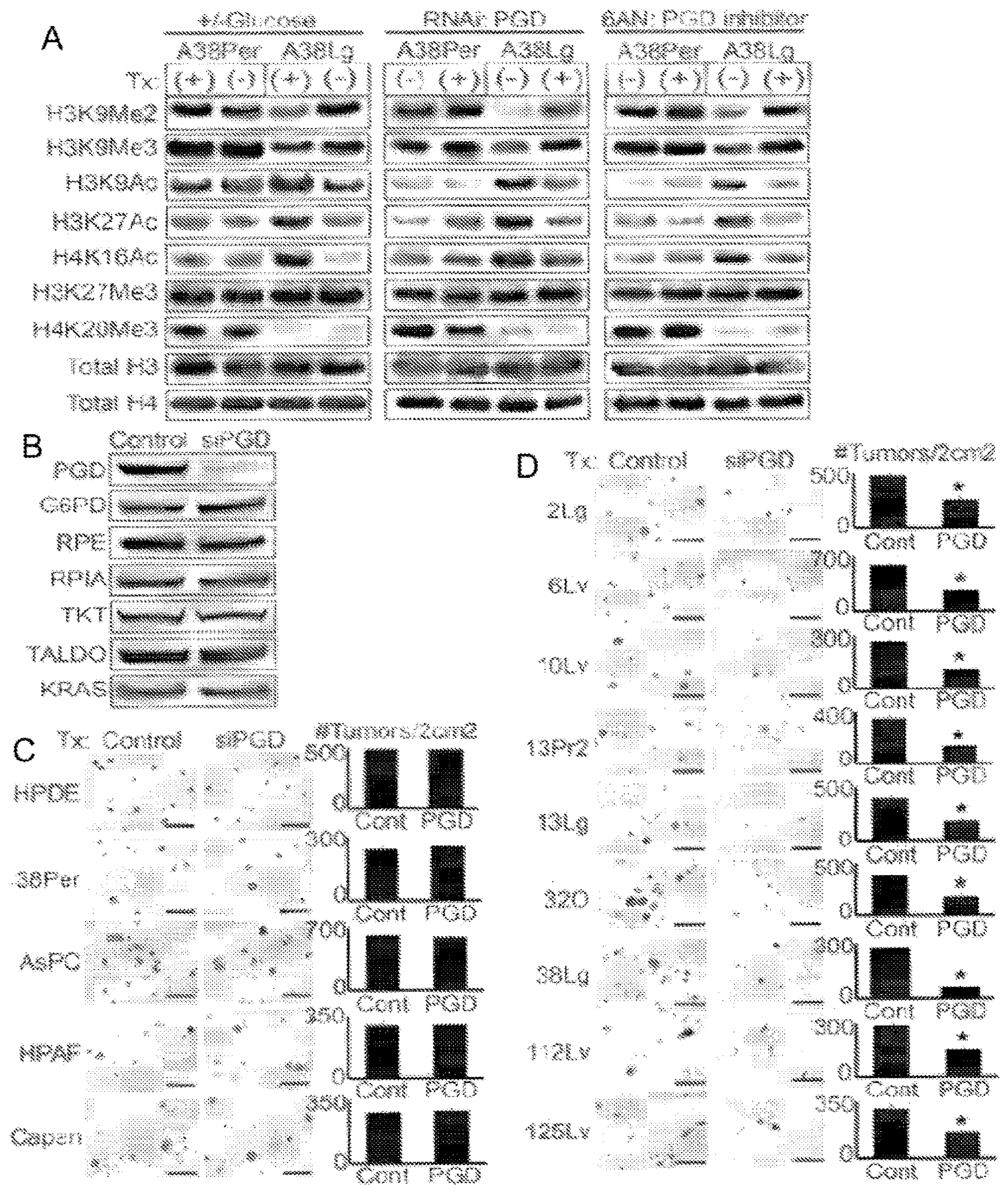


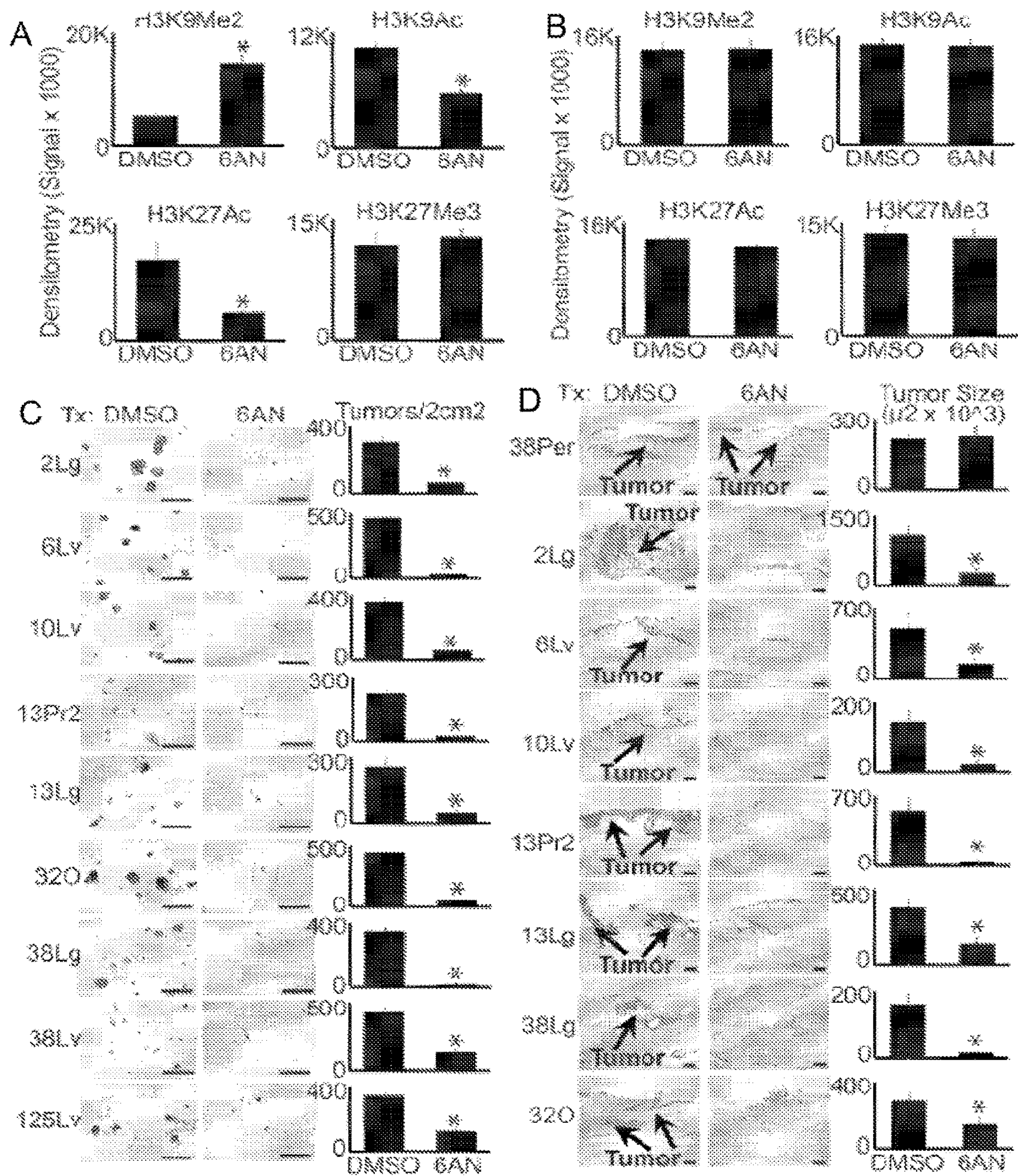
FIGURE 3G



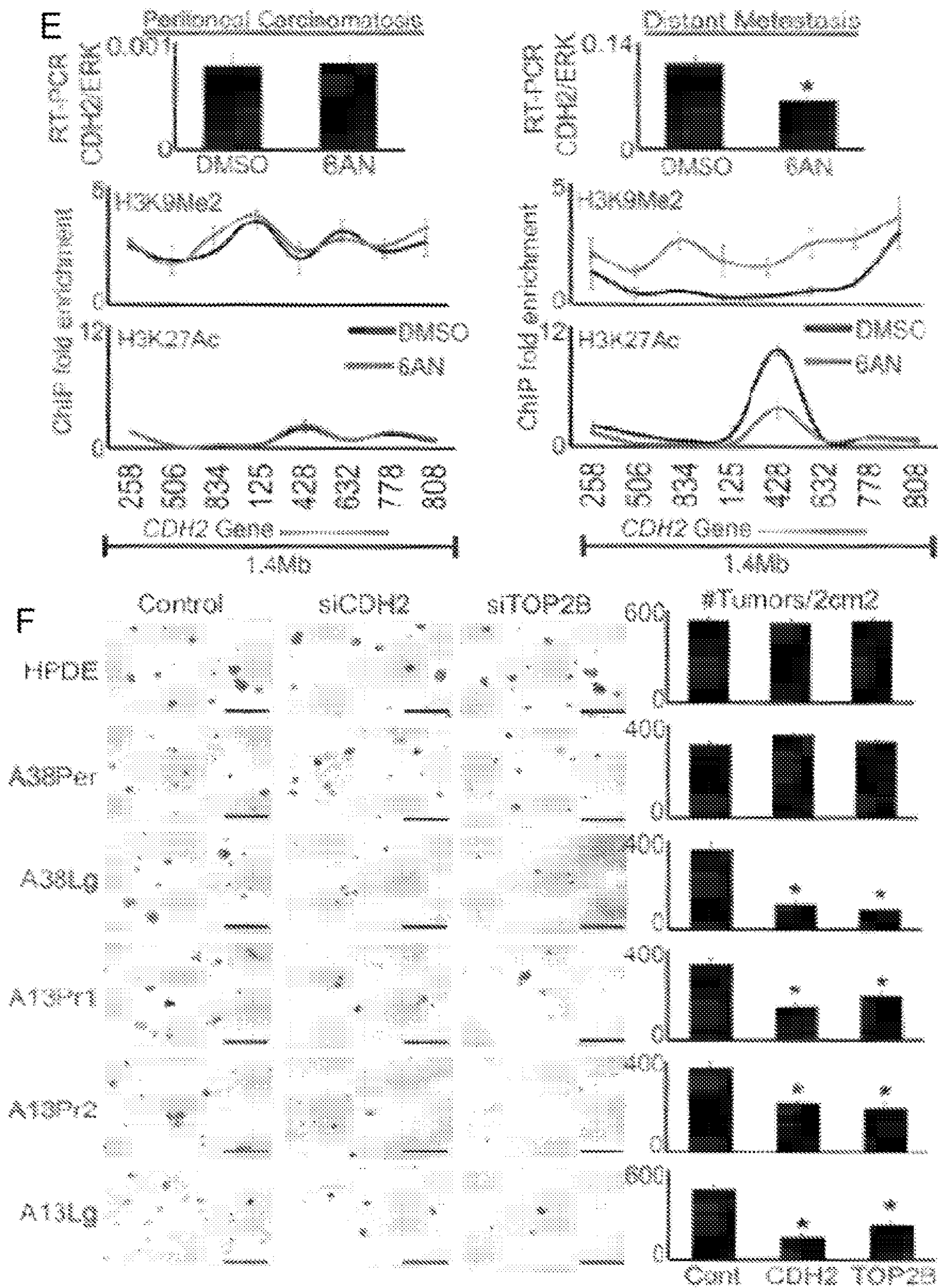
FIGURES 4A-4F



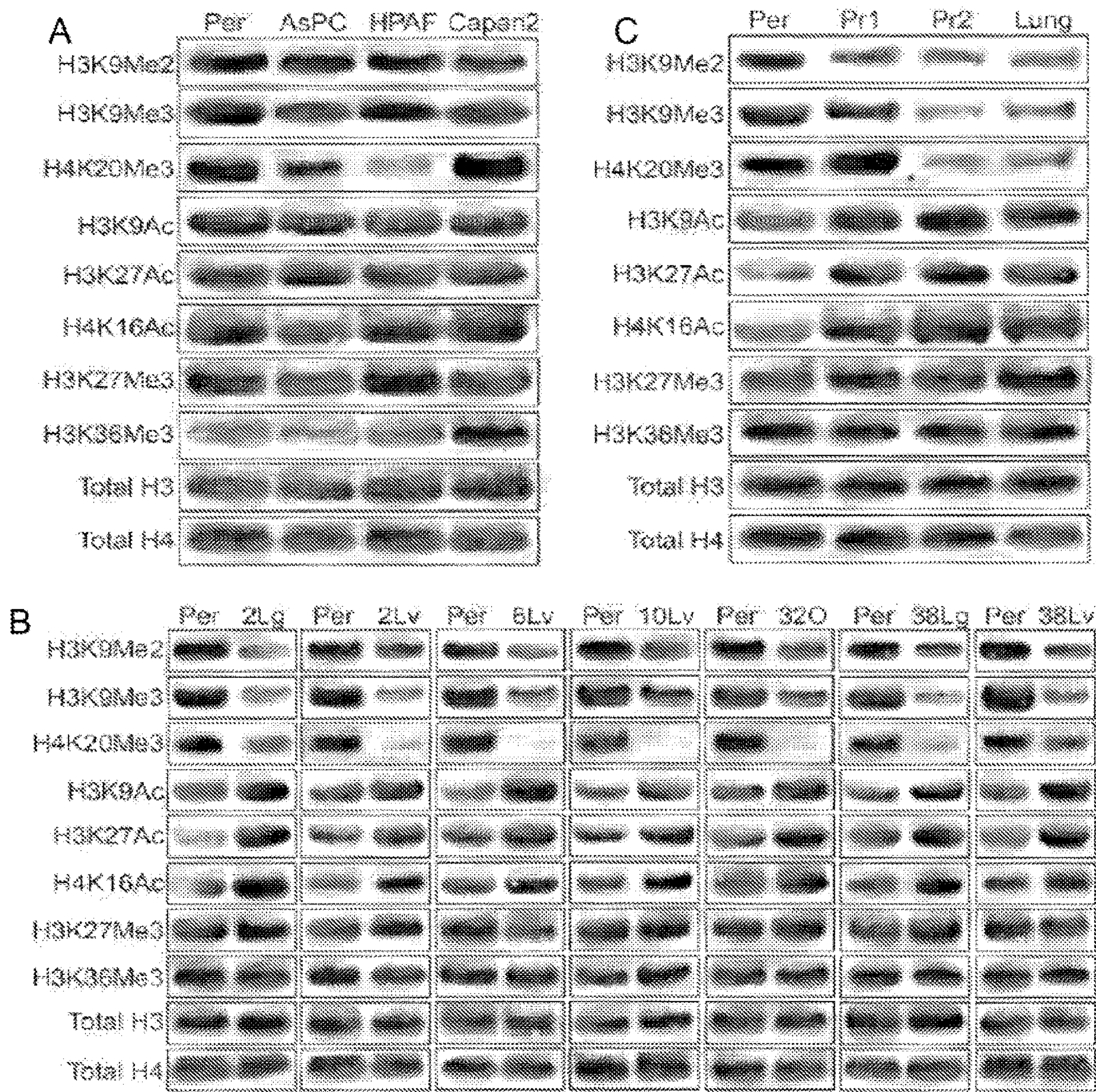
FIGURES 5A-5D



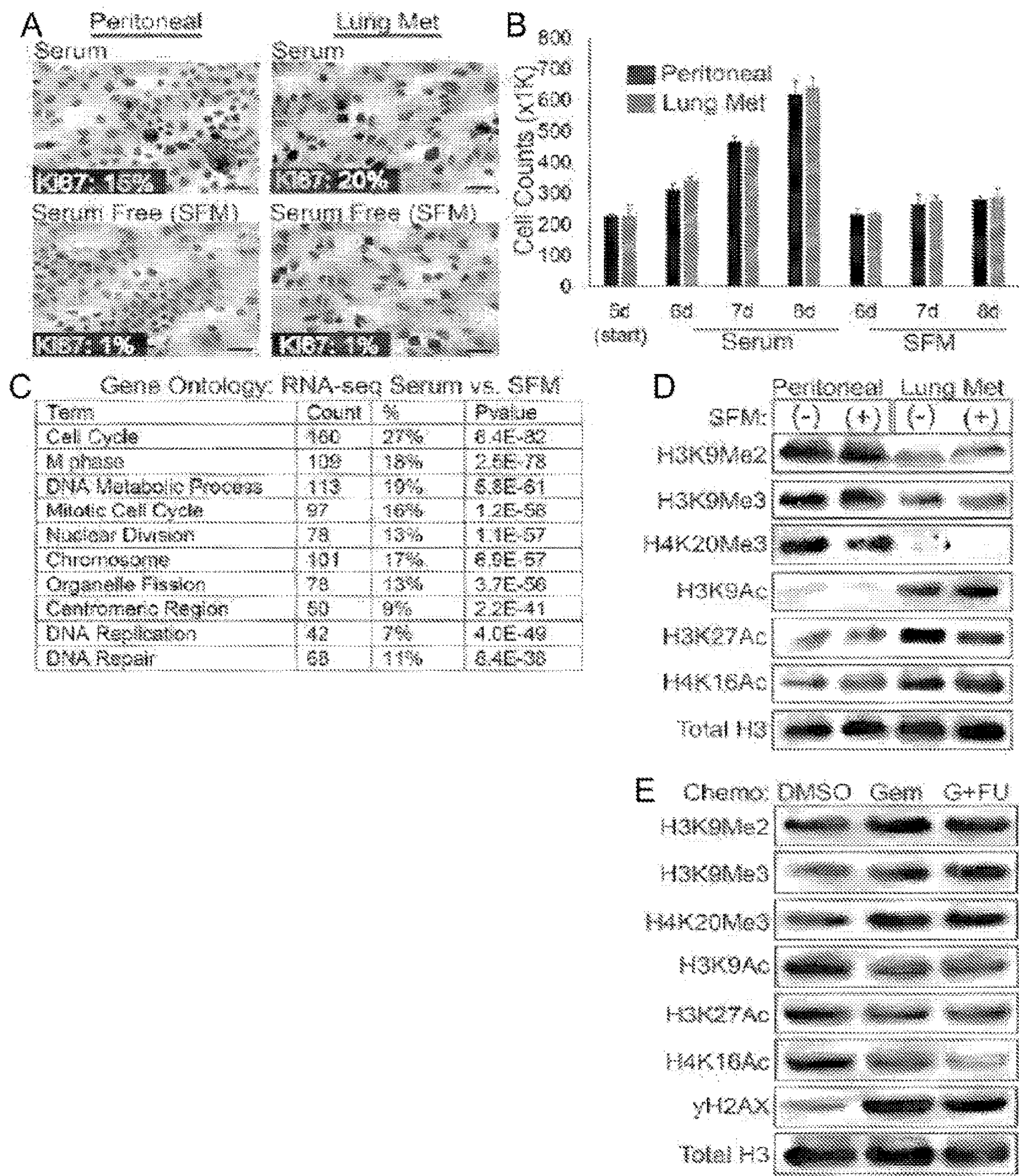
FIGURES 6A-6D



FIGURES 6E-6F



FIGURES 7A-7C



FIGURES 8A-8E

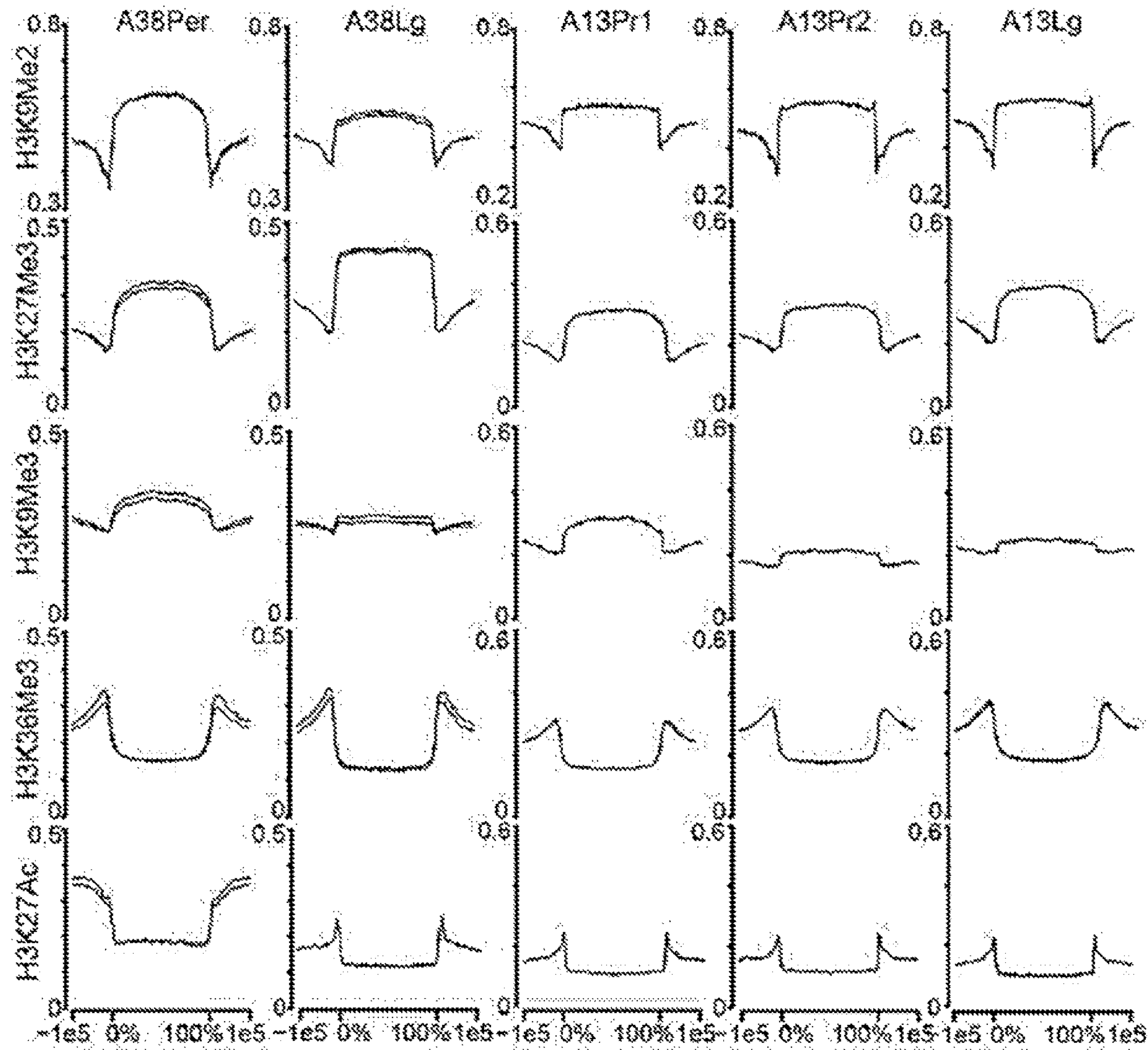
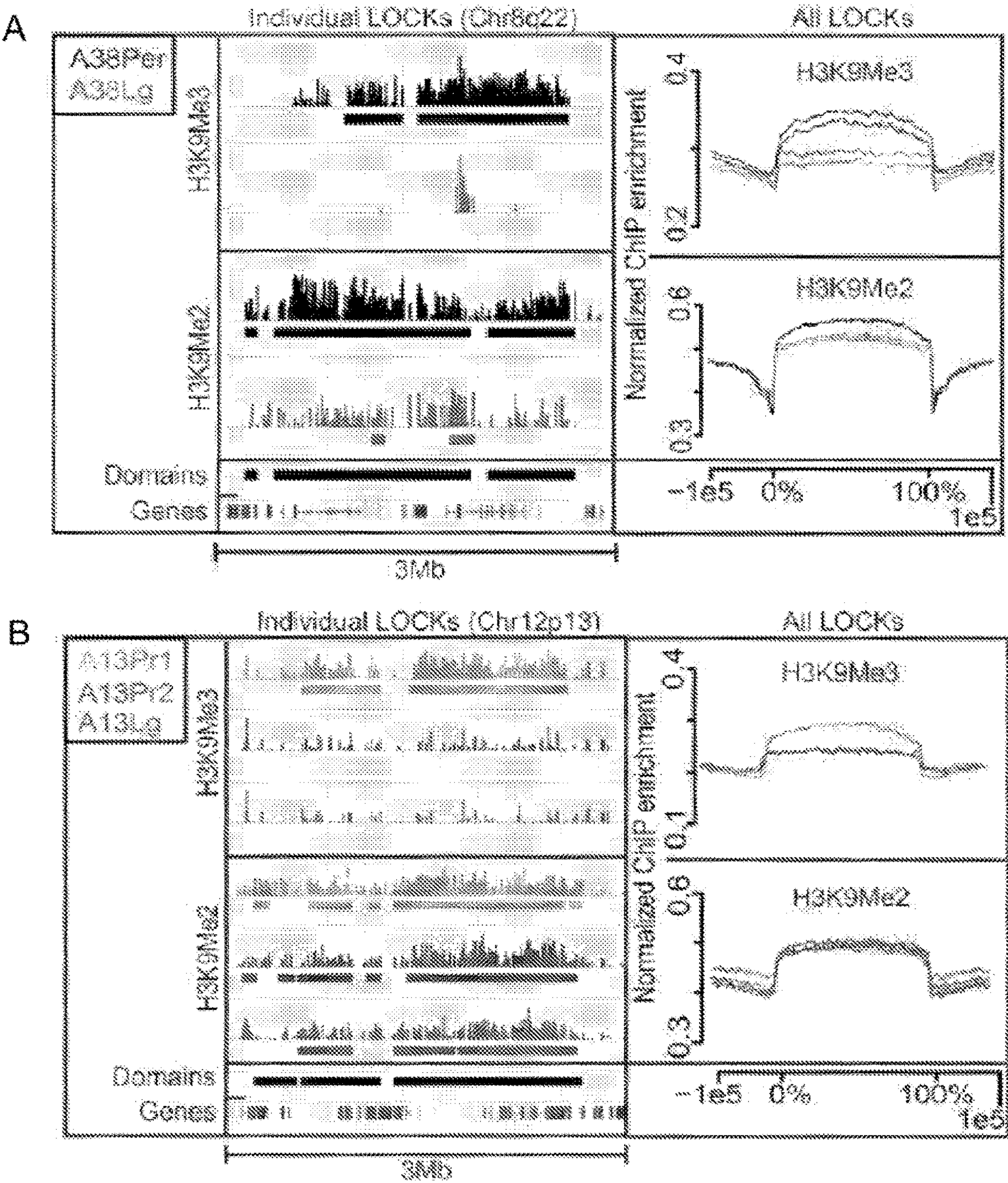
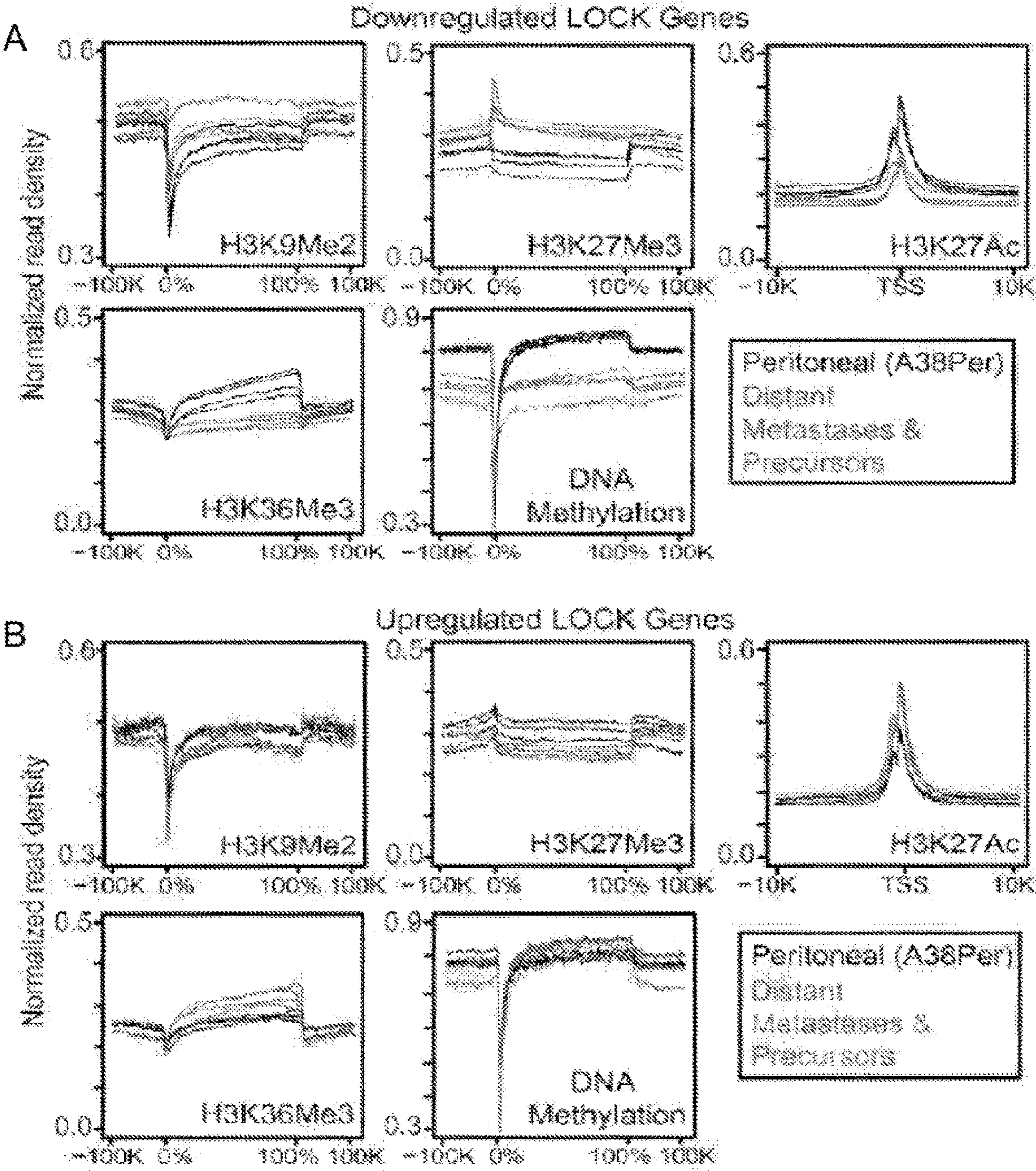


FIGURE 9



FIGURES 10A-10B



FIGURES 11A-11B

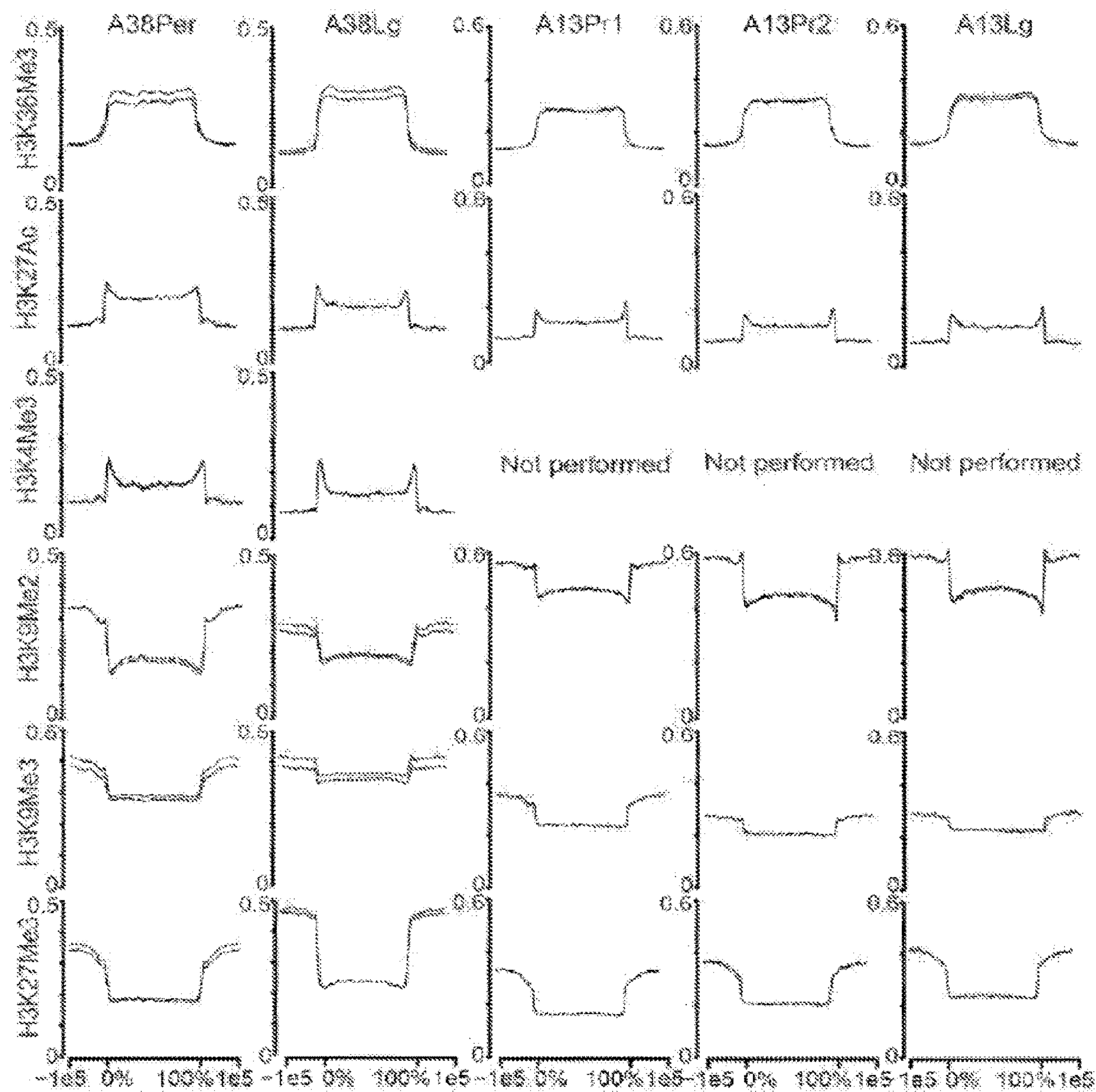
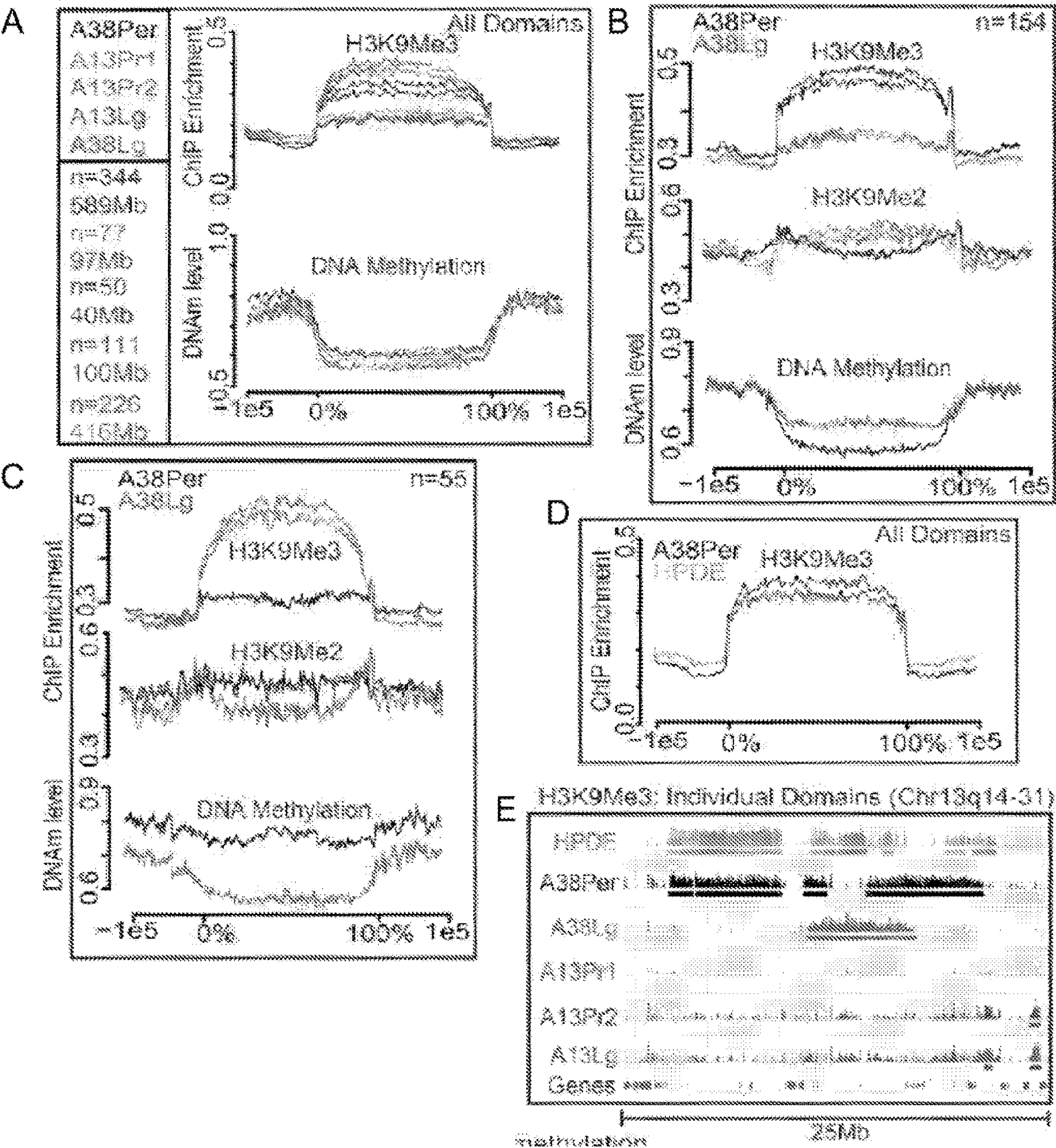


FIGURE 12



FIGURES 13A-13E

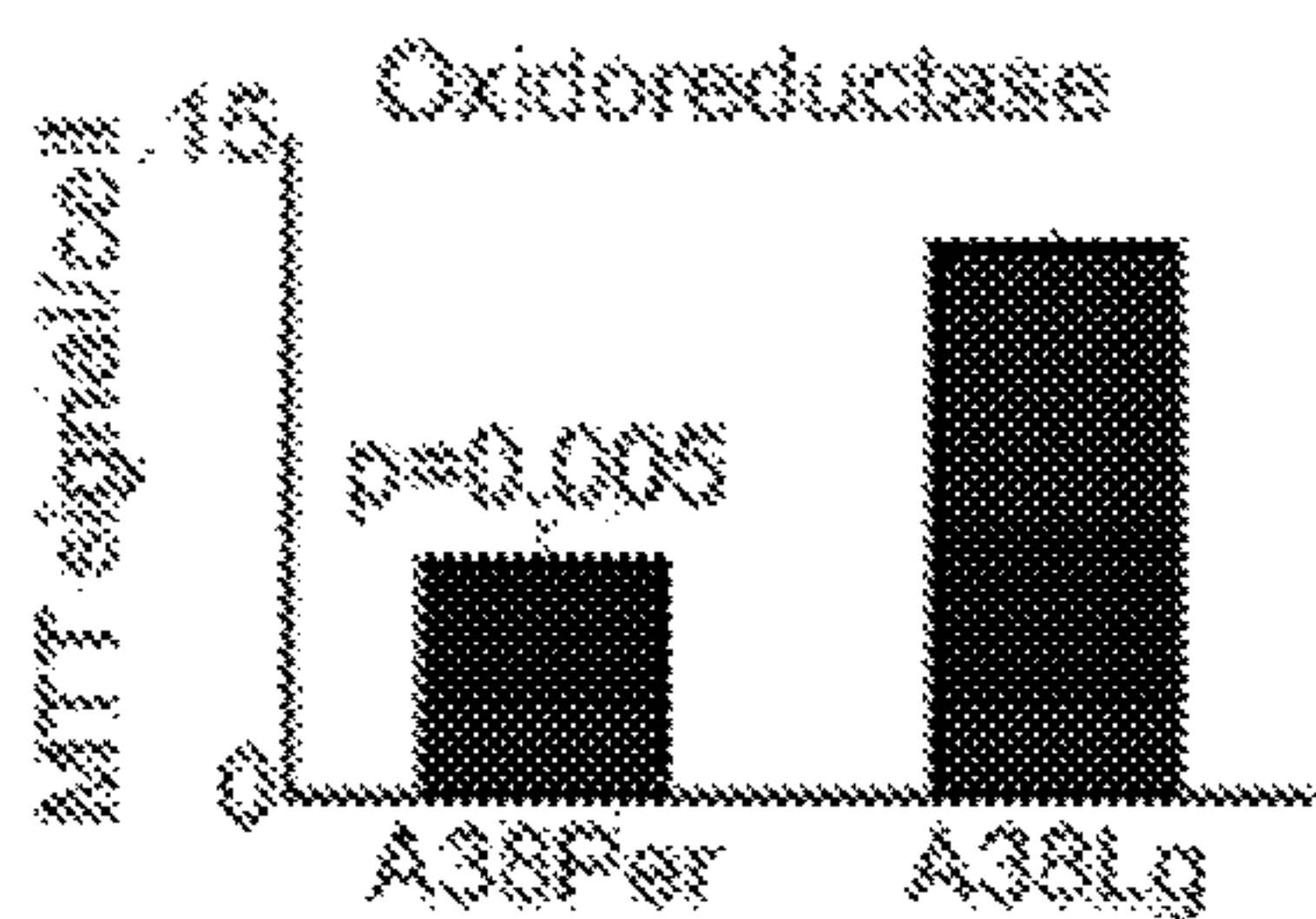


FIGURE 14A

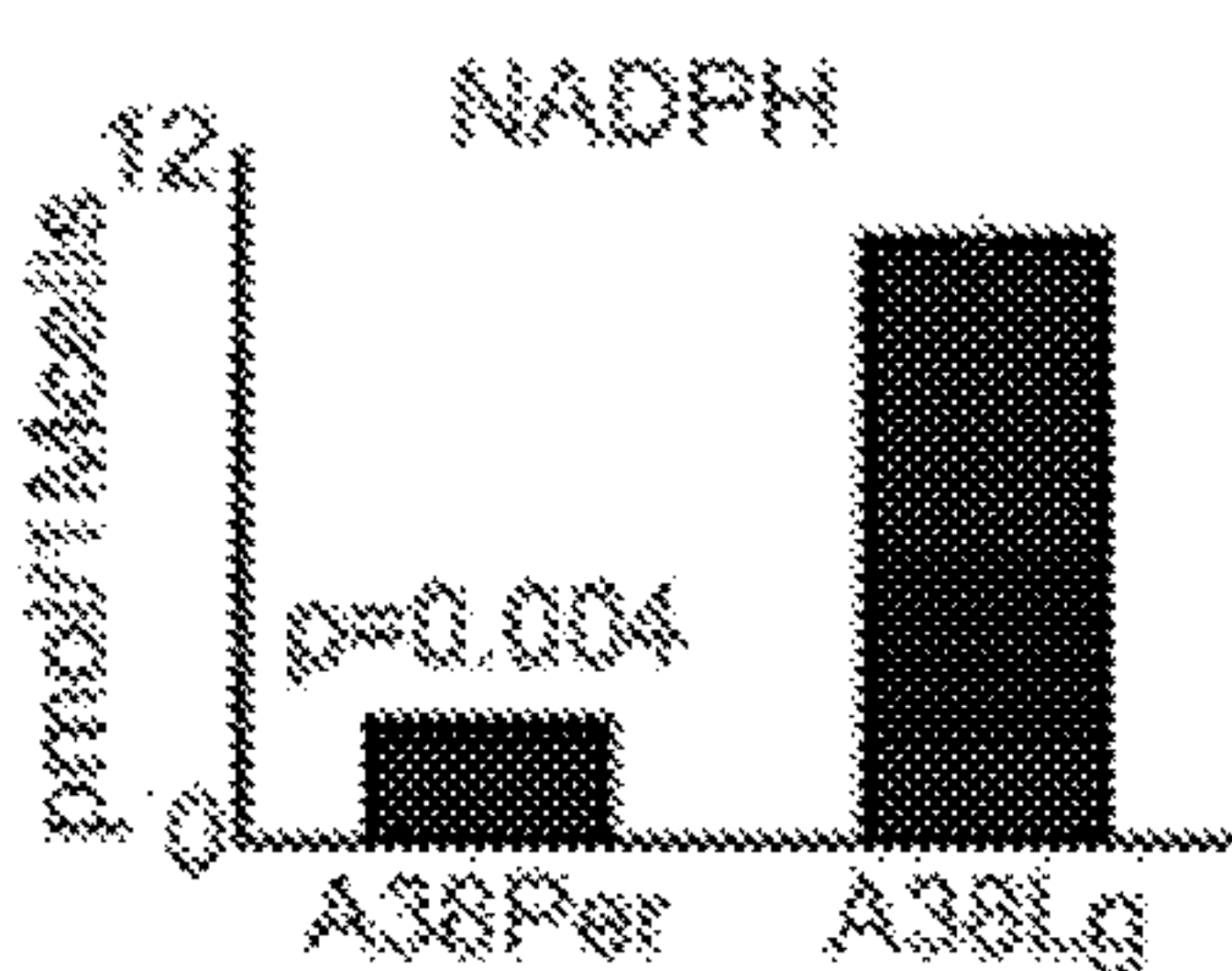


FIGURE 14B

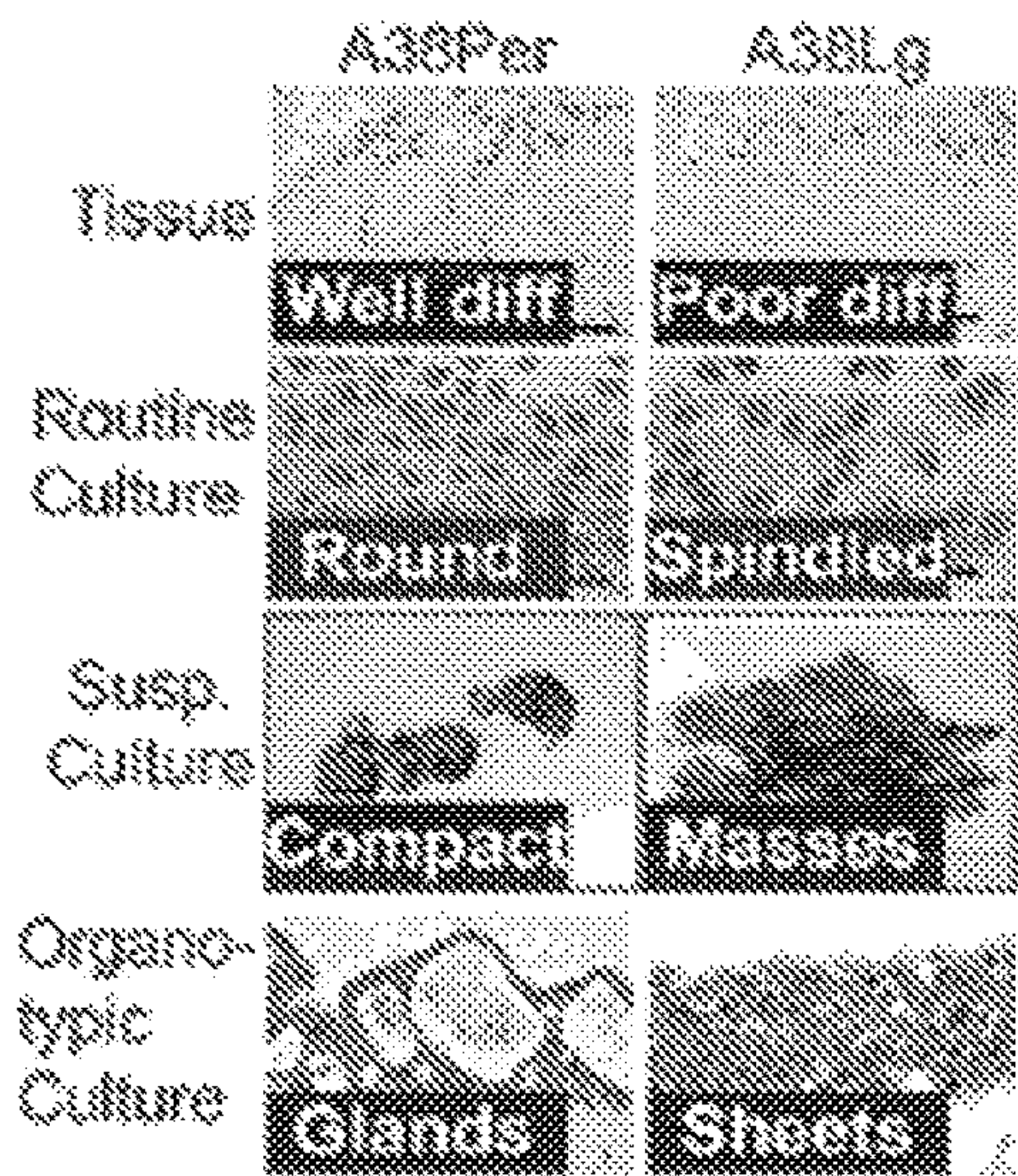
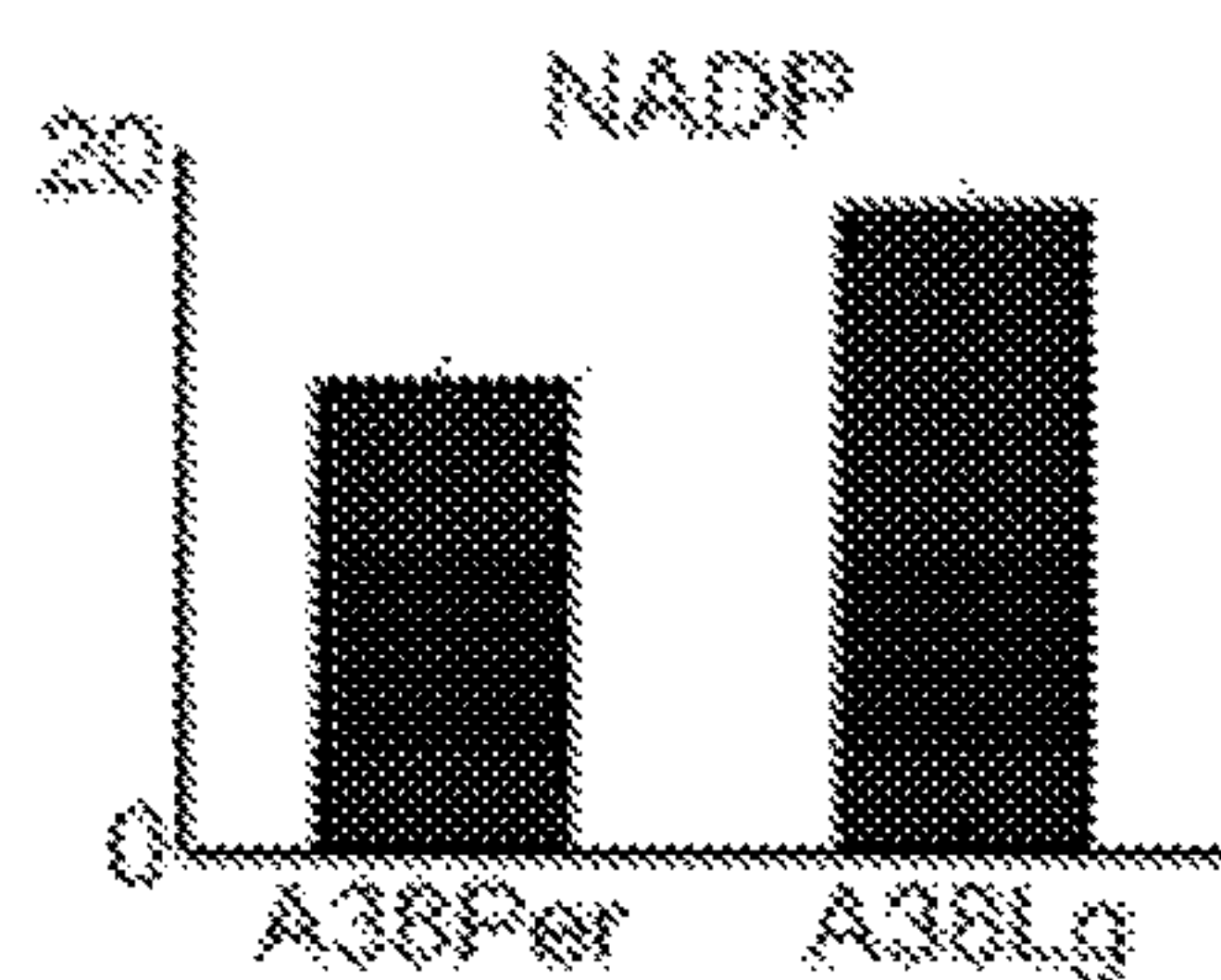


FIGURE 14C

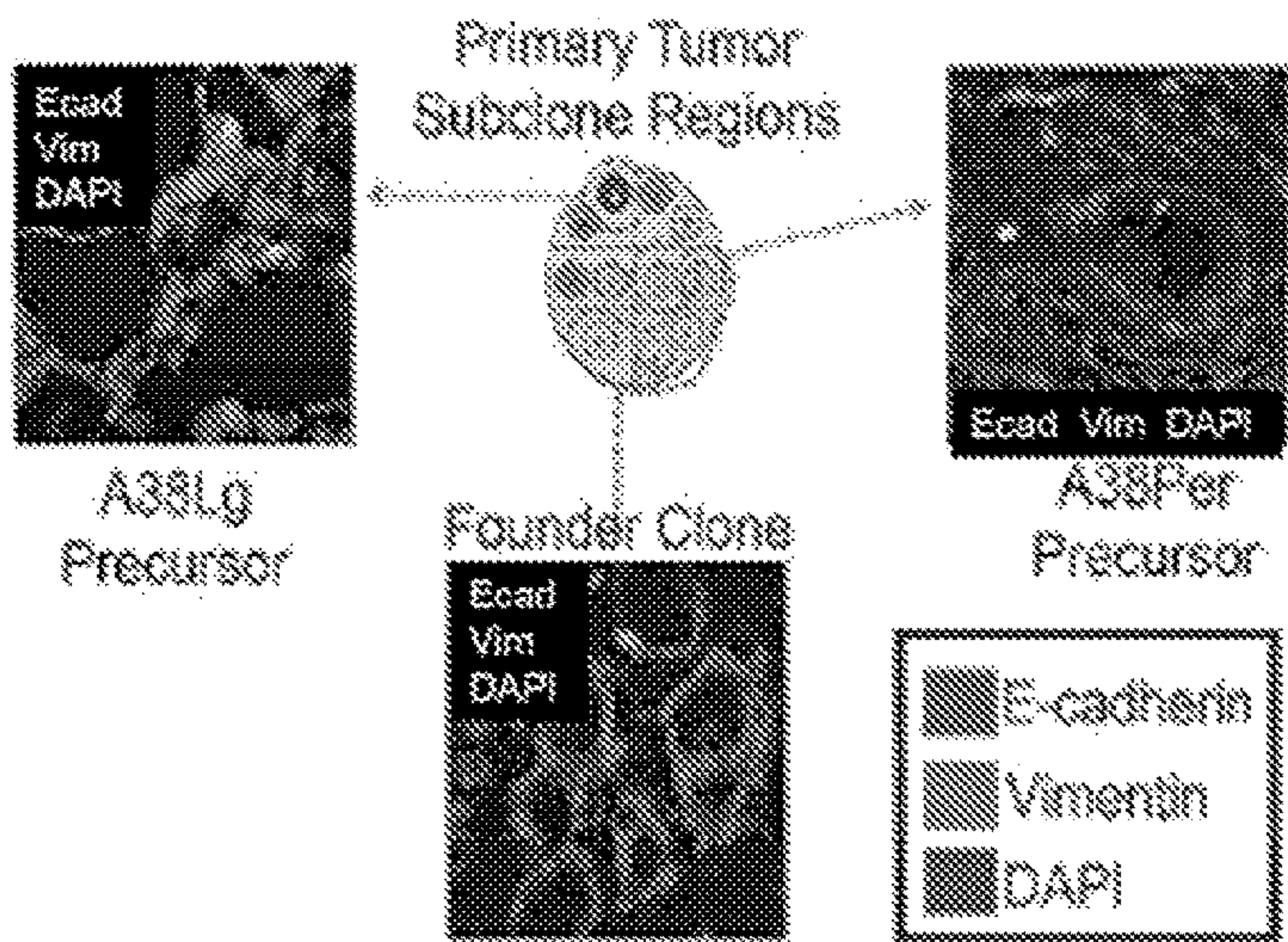


FIGURE 14D

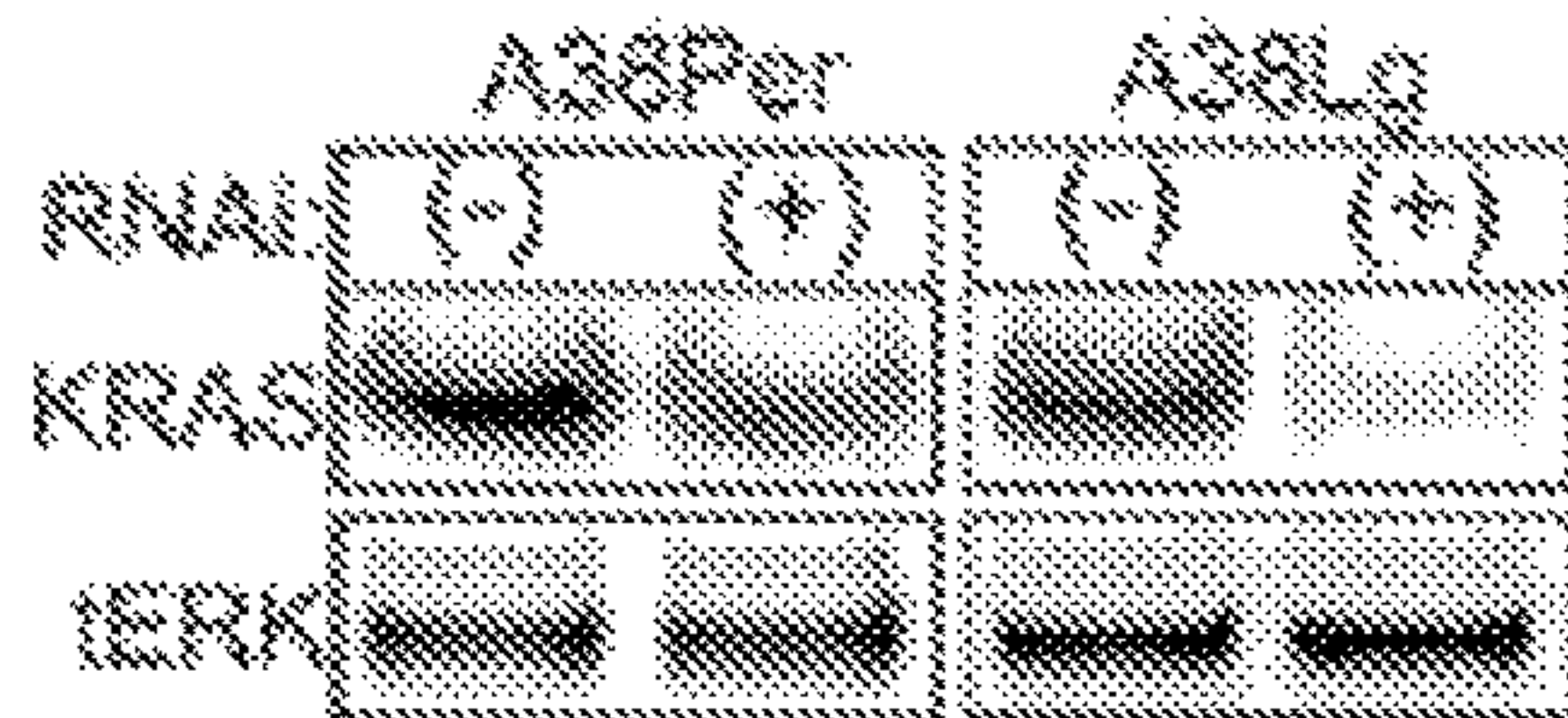


FIGURE 14E

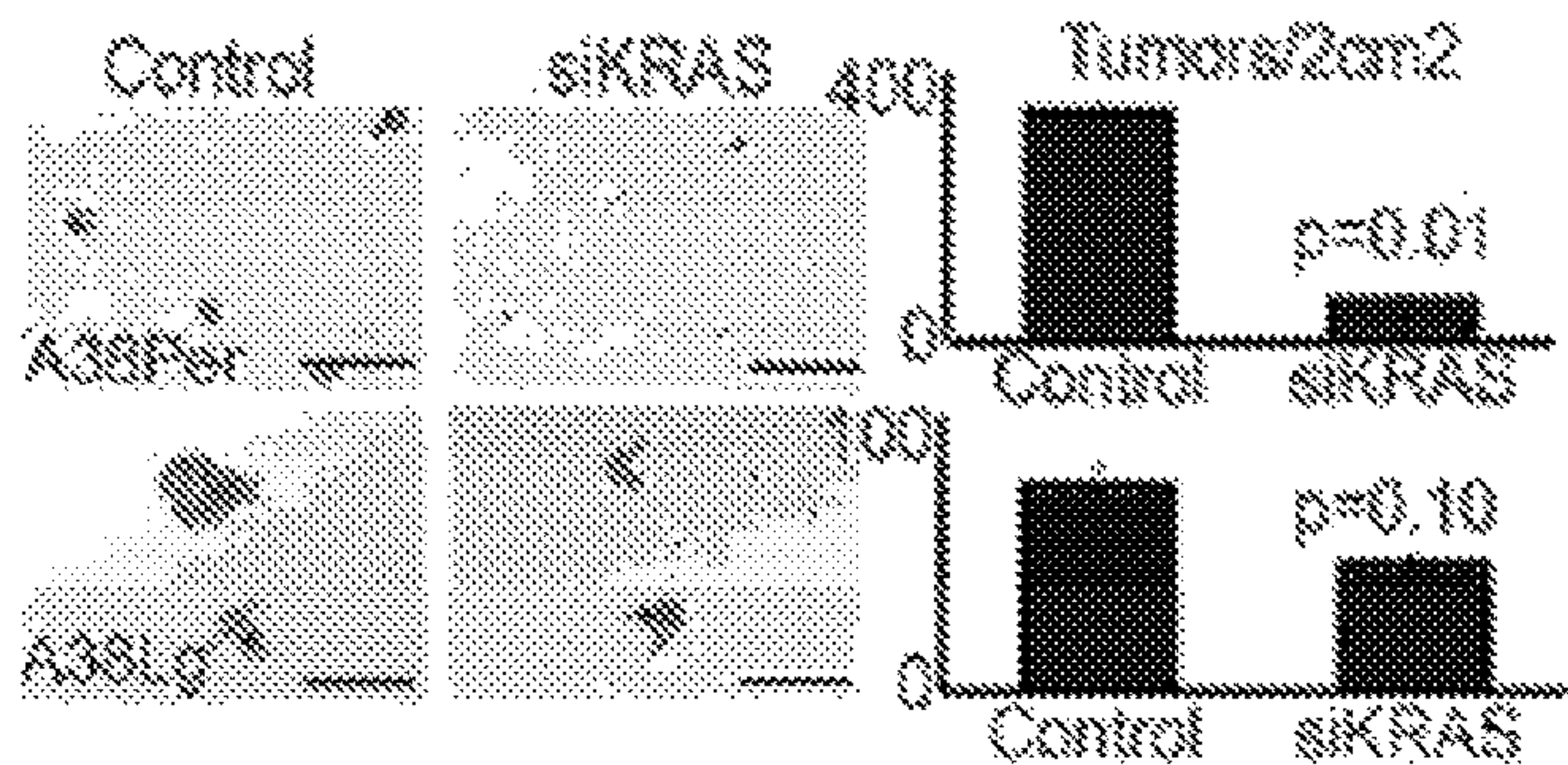
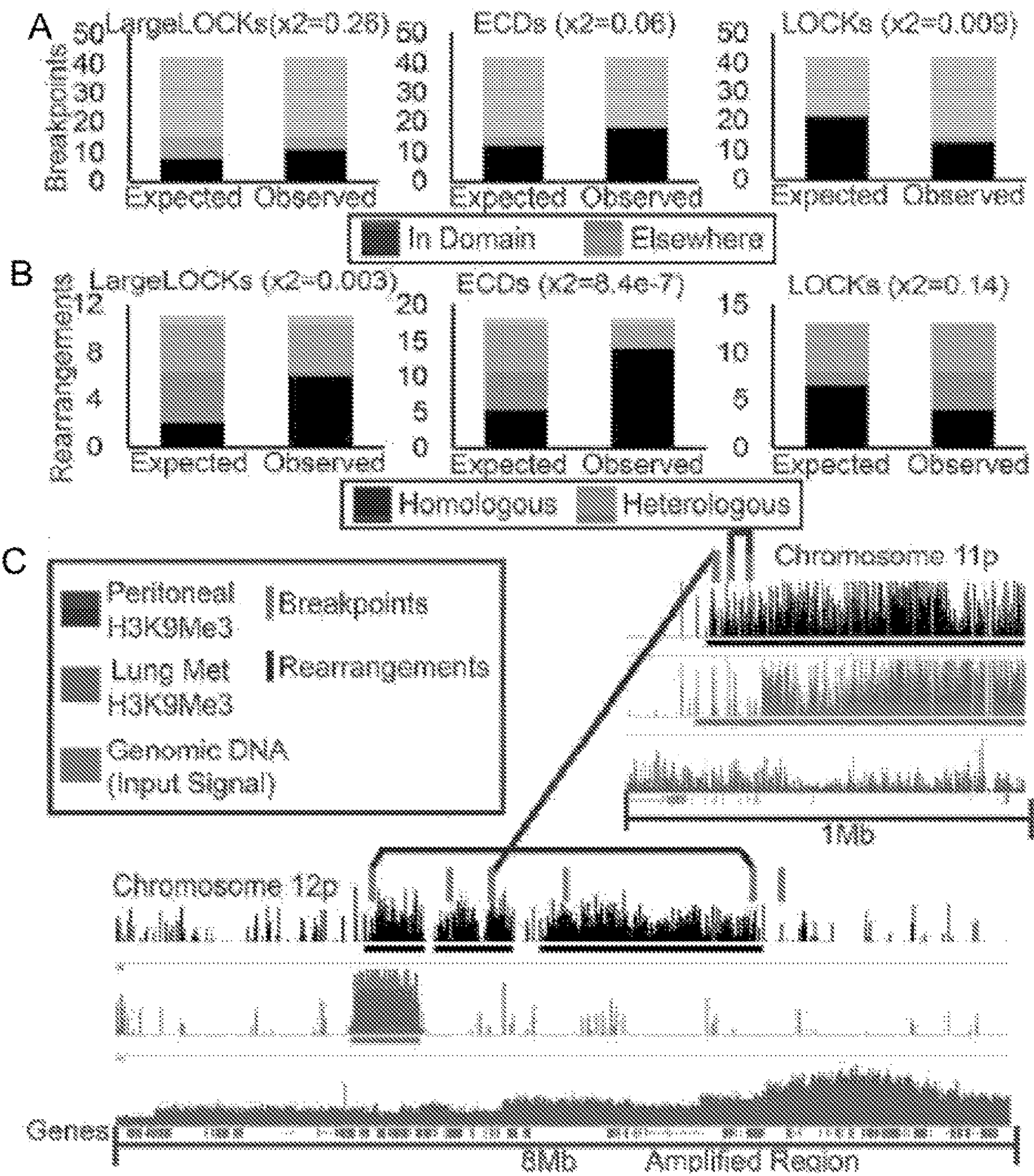
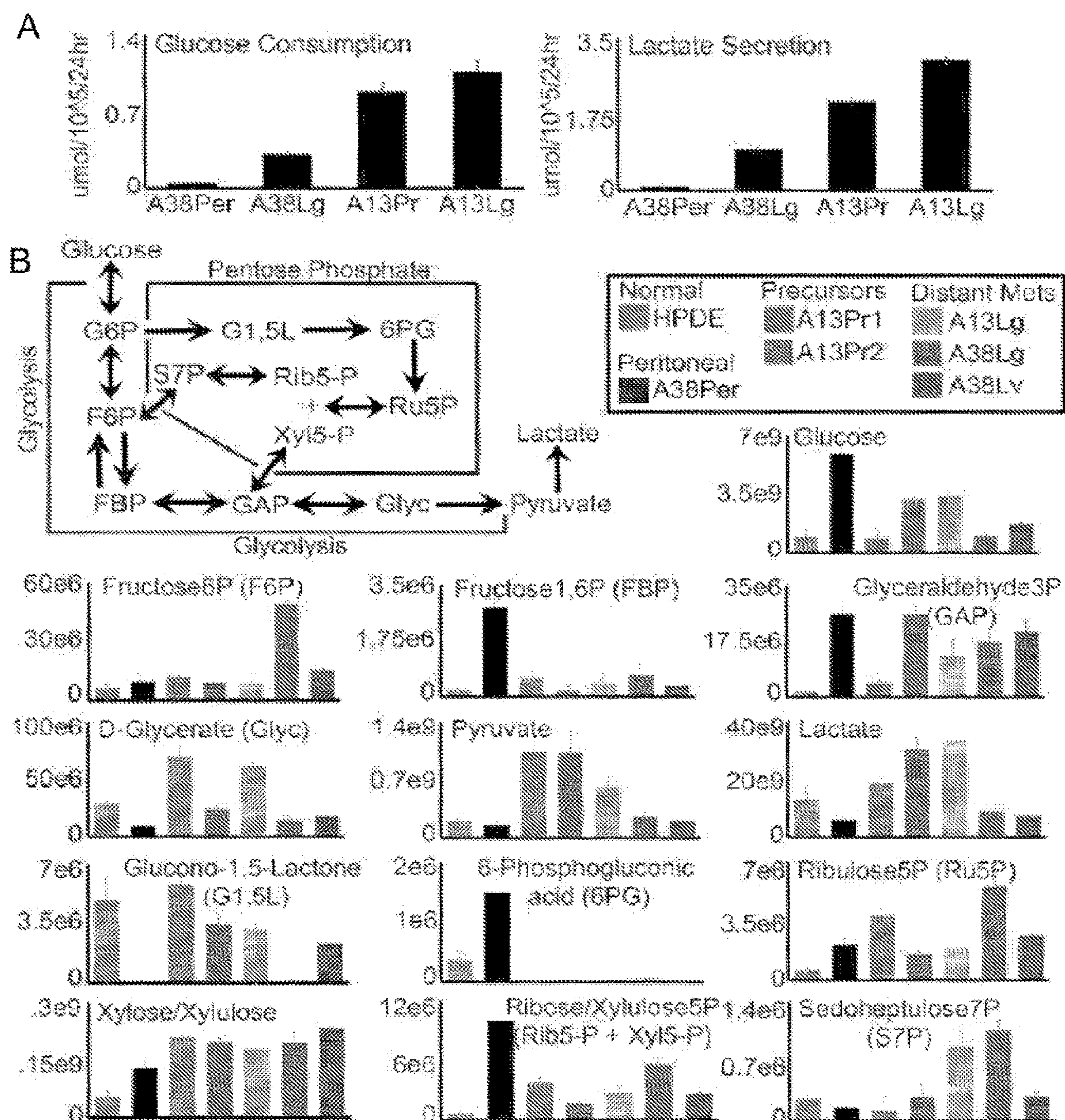


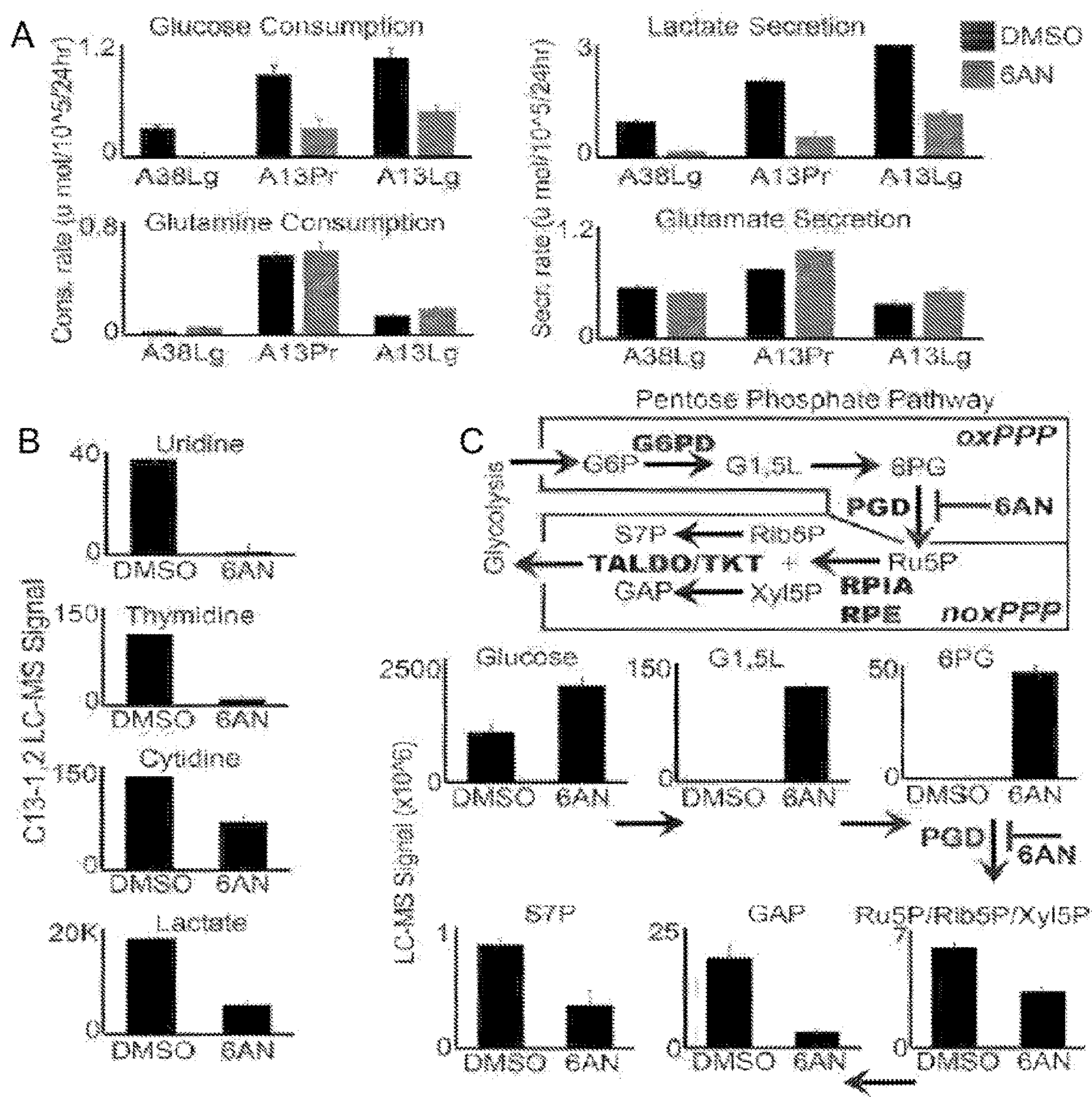
FIGURE 14F



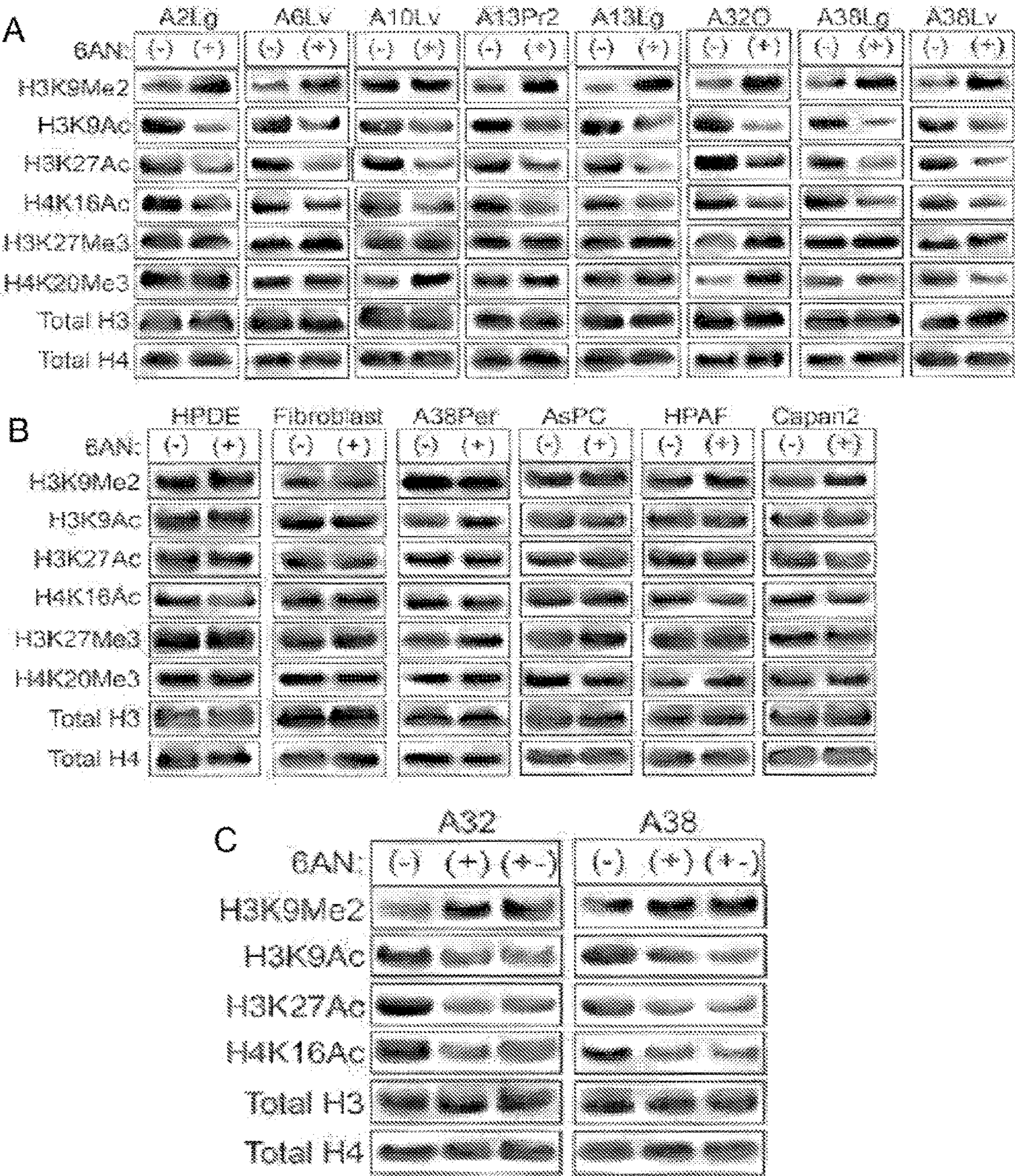
FIGURES 15A-15C



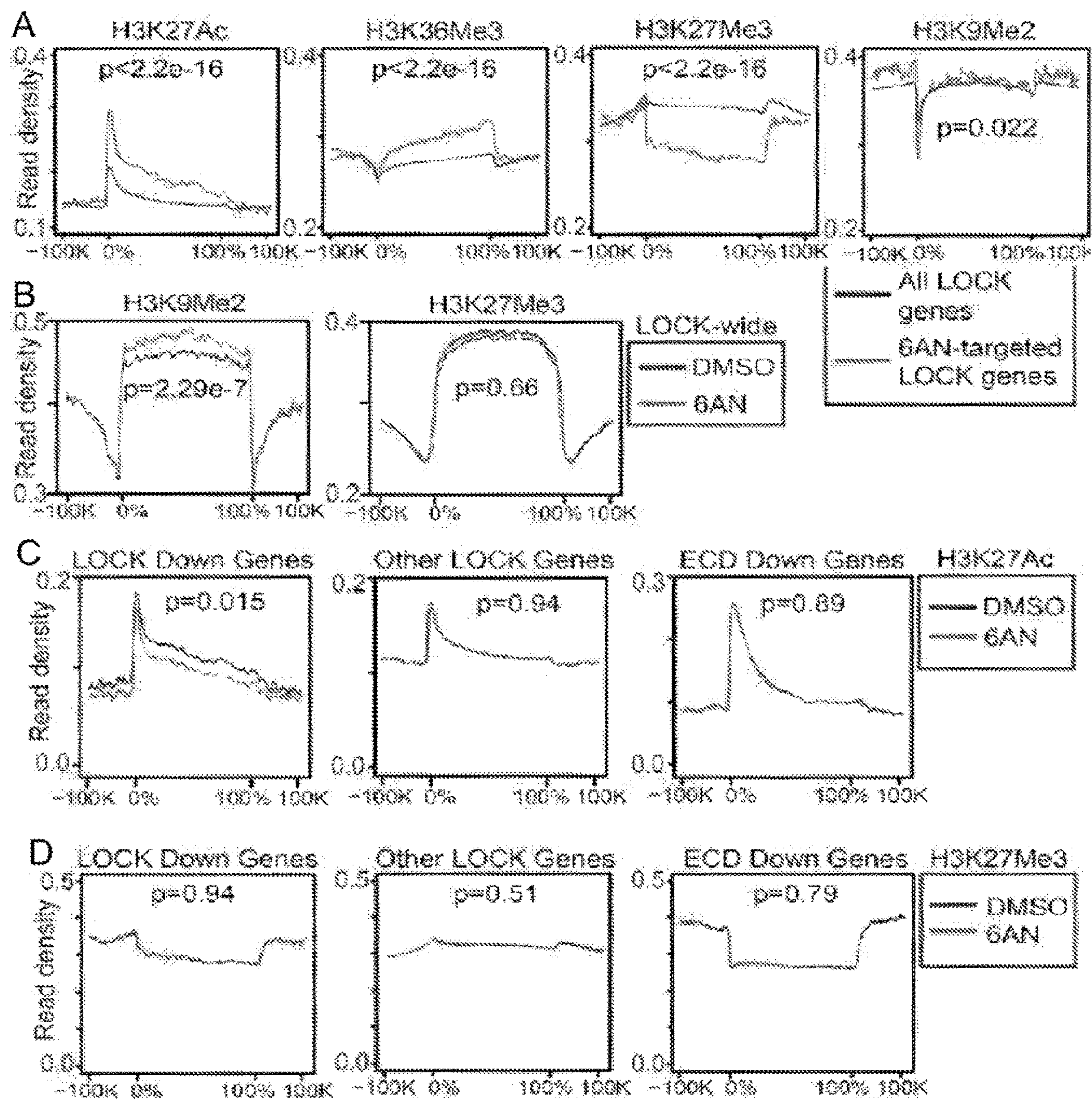
FIGURES 16A-16B



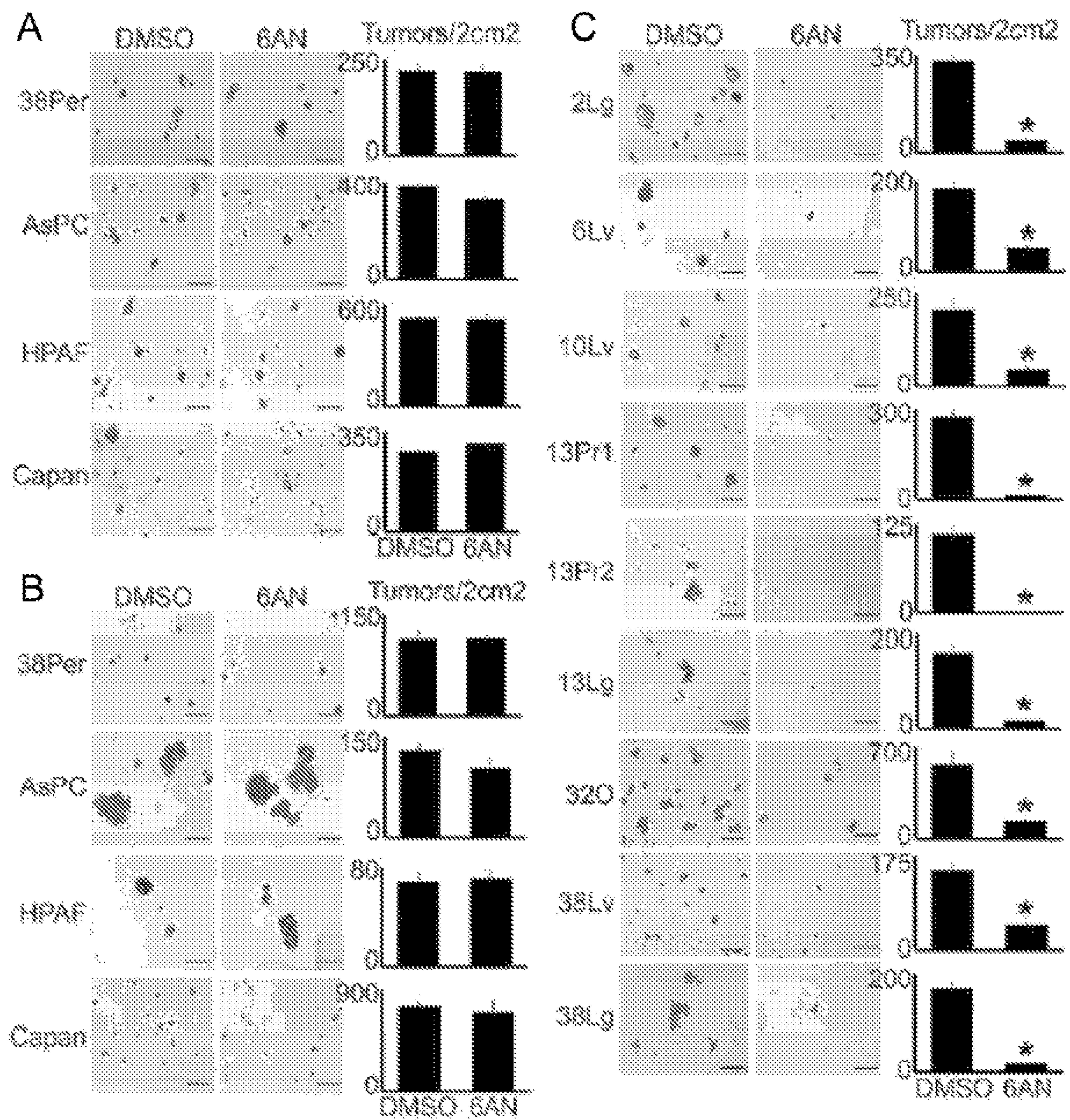
FIGURES 17A-17C



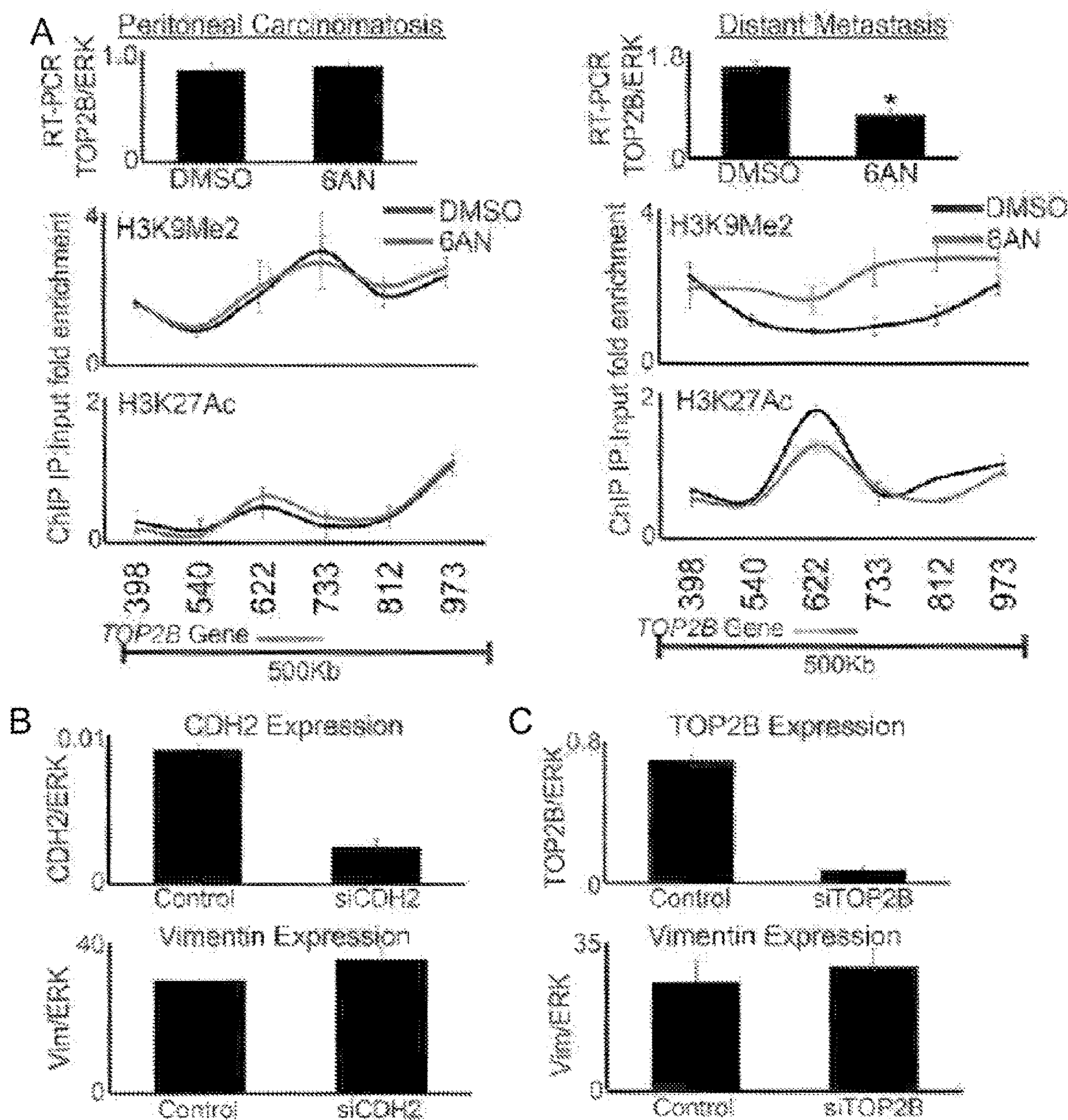
FIGURES 18A-18C



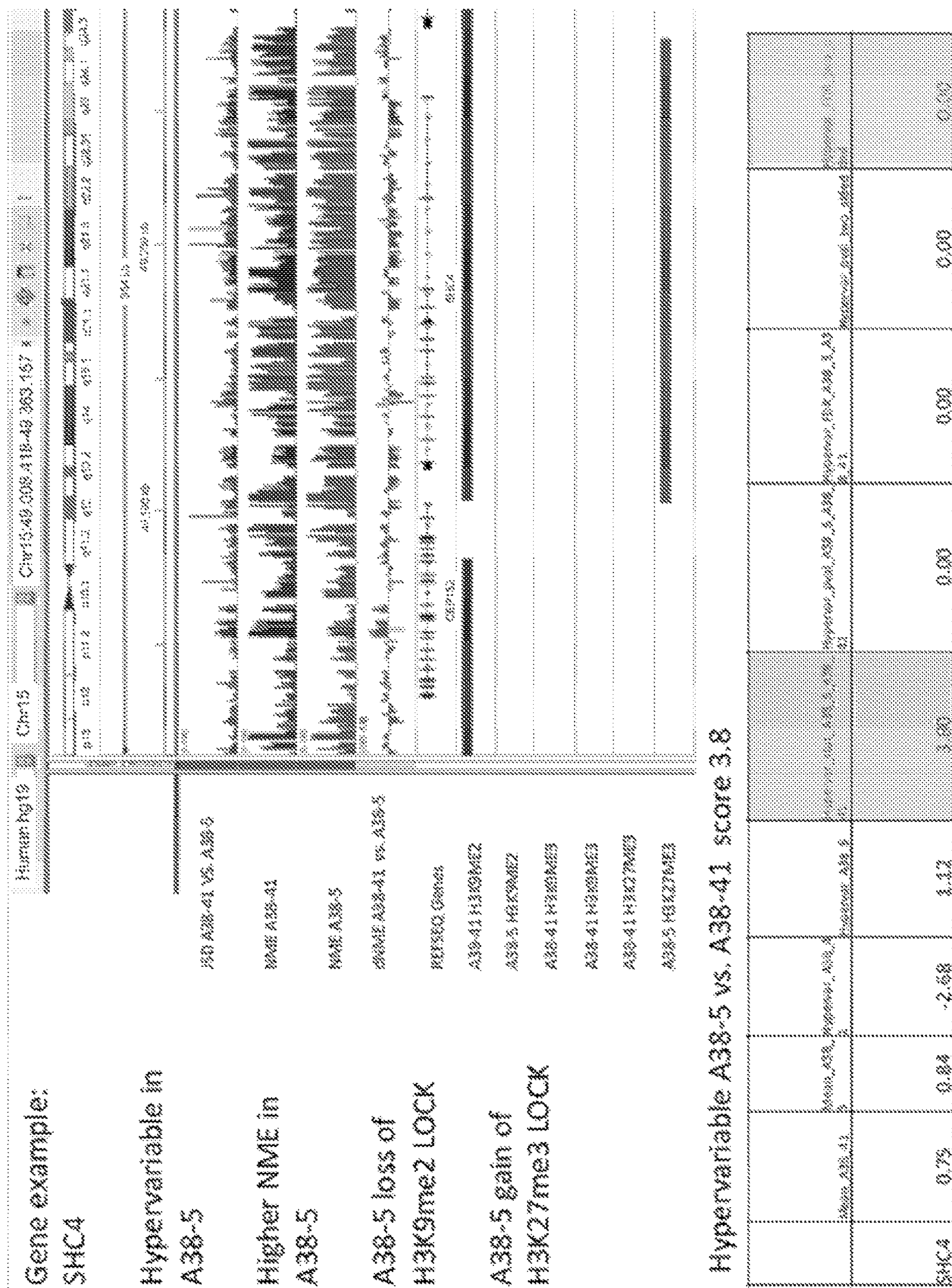
FIGURES 19A-19D



FIGURES 20A-20C



FIGURES 21A-21C



LARGE-SCALE EPIGENOMIC REPROGRAMMING LINKS ANABOLIC GLUCOSE METABOLISM TO DISTANT METASTASIS DURING THE EVOLUTION OF PANCREATIC CANCER PROGRESSION

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application is a divisional application of U.S. application Ser. No. 16/340,069 filed Apr. 5, 2019, now pending; which is a 35 USC § 371 National Stage application of International Application No. PCT/US2017/055376 filed Oct. 5, 2017, now expired; which claims the benefit under 35 USC § 119(e) to U.S. Application Ser. No. 62/405,155 filed Oct. 6, 2016, now expired. The disclosure of each of the prior applications is considered part of and is incorporated by reference in the disclosure of this application.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

[0002] This invention was made with government support under CA054358, CA140599, CA179991, CA180682 and CA095103 awarded by the National Institutes of Health. The government has certain rights in the invention.

INCORPORATION OF SEQUENCE LISTING

[0003] The material in the accompanying sequence listing is hereby incorporated by reference into this application. The accompanying sequence listing .xml file, named JHU4080-2_ST26.xml, was created on Jul. 19, 2023 and is 55,580 bytes in size.

BACKGROUND OF THE INVENTION

Field of the Invention

[0004] The invention relates generally to genetic analysis and more specifically to cancer and the epigenetic influence on progression and metastases of cancer.

Background Information

[0005] During the evolutionary progression of pancreatic ductal adenocarcinoma (PDAC), heterogeneous subclonal populations emerge that drive primary tumor growth, regional spread, distant metastasis, and patient death. However, the genetics of metastases largely reflects that of the primary tumor in untreated patients, and PDAC driver mutations are shared by all subclones. This raises the possibility that an epigenetic process might be operative during metastasis. Here we detected striking epigenetic reprogramming of global chromatin modifications during the natural evolutionary history of distant metastasis. Genome-wide mapping revealed that these global changes were targeted to thousands of large chromatin domains across the genome that collectively specified malignant traits, including euchromatin and large organized chromatin K9-modified (LOCK) heterochromatin. Parallel to these changes, distant metastases co-evolved a dependence on the oxidative branch of the pentose phosphate pathway (oxPPP), and oxPPP inhibition selectively reversed reprogrammed chromatin and blocked tumorigenic potential. Thus, diver-

gent metabolic, epigenetic, and tumorigenic programs emerged during the evolution of pancreatic cancer progression.

[0006] Despite significant progress in survival rates for most human cancers, PDAC remains nearly universally lethal with survival rates of 8%. In fact, PDAC is projected to be the second-leading cause of cancer deaths in the western world by 2020. Primary PDACs have been shown to contain distinct subclonal populations. However, these subclones share identical driver mutations and the genetics of metastases largely reflects that of the primary tumor. Furthermore, subclones are defined genetically by their unique progressor mutations, the vast majority if not all of which are thought to be passenger events. This raises questions as to what mechanisms might drive progression and metastasis during the natural history of disease evolution.

[0007] One prometastatic candidate is epigenomic regulation. In particular, the inventors wished to investigate the role of large-scale epigenomic changes during PDAC subclonal evolution and distant metastasis, especially within heterochromatin domains including large organized heterochromatin lysine (K)-9 modified domains (LOCKs) and large DNA hypomethylated blocks. These regions could represent selectable targets for large-scale epigenetic reprogramming, since they occupy over half of the genome, partially overlap with one another, and are found in many human cancers including PDAC. It was therefore hypothesized that epigenomic dysregulation within these regions could be a major selective force for tumor progression, given the lack of any consistent metastasis-specific driver mutations.

SUMMARY OF THE INVENTION

[0008] The present invention relates to a method of identifying targets for epigenetic reprogramming comprising detecting large organized heterochromatin lysine (K)-9 modified domains (LOCKs) and large DNA hypomethylated blocks in a sample containing DNA from a subject having cancer. For example, the method applies to a subject that has or is at risk of having PDAC and/or metastasis thereof. In one aspect, the detection comprises analysis of H3K9Me2/3 and/or H4K20Me3. In another aspect, the detection comprises analysis of H3K27Ac and/or H3K9Ac.

[0009] In another embodiment, the invention provides for the use of differentially expressed genes to identify metastatic propensity in primary tumors, wherein the genes are selected from genes in the Tables herein, oxidative stress genes, EMT genes, immunological response genes, DNA repair genes, glucose metabolism genes, oxPPP genes, and PGD genes.

[0010] In another embodiment, the invention provides a method for identifying agents or compounds to affect epigenomic changes, including inhibition of oxPPP comprising analyzing a sample from a subject before and after contacting with the agent or compound and determining the effect of the agent or compound on the epigenomic changes.

BRIEF DESCRIPTION OF THE DRAWINGS

[0011] FIGS. 1A-1E relate to global epigenetic reprogramming during the evolution of distant metastasis.

[0012] FIG. 1A is a series of immunohistochemical stains.

[0013] FIG. 1B is a series of immunohistochemical stains.

- [0014] FIG. 1C is a series of immunohistochemical stains.
- [0015] FIG. 1D is a series of immunohistochemical stains.
- [0016] FIG. 1E is a series of immunohistochemical stains.
- [0017] FIG. 1F is a series of western blot images.
- [0018] FIG. 1G is a series of graphical representations of data.
- [0019] FIGS. 2A-2E relate to epigenomic reprogramming of chromatin domains during PDAC subclonal evolution.
- [0020] FIG. 2A is a graphical representation of data.
- [0021] FIG. 2B is a graphical representation of data.
- [0022] FIG. 2C is a graphical representation of data.
- [0023] FIG. 2D is a graphical representation of data.
- [0024] FIG. 2E is a graphical representation of data.
- [0025] FIGS. 3A-3G relate to reprogrammed chromatin domains encoding divergent malignant properties.
- [0026] FIG. 3A is a graphical representation of data.
- [0027] FIG. 3B is a series of western blot images.
- [0028] FIG. 3C is a graphical representation of data.
- [0029] FIG. 3D is a series of western blot images.
- [0030] FIG. 3E is a series of western blot images.
- [0031] FIG. 3F is a graphical representation of data.
- [0032] FIG. 3G is a series of images of tumor forming assays and related graphical plots.
- [0033] FIGS. 4A-4F relate to hyperactive glucose metabolism and 6PG depletion in distant metastatic subclones.
- [0034] FIG. 4A is a graphical representation of data.
- [0035] FIG. 4B is a graphical representation of data.
- [0036] FIG. 4C is a graphical representation of data.
- [0037] FIG. 4D is a schematic diagram.
- [0038] FIG. 4E is a series of graphical representations of data.
- [0039] FIG. 4F is a graphical representation of data.
- [0040] FIGS. 5A-5D relate to PGD-dependence in distant metastatic subclones.
- [0041] FIG. 5A is a series of western blot images.
- [0042] FIG. 5B is a series of western blot images.
- [0043] FIG. 5C is a series of images of tumor forming assays and related graphical plots.
- [0044] FIG. 5D is a series of images of tumor forming assays and related graphical plots.
- [0045] FIGS. 6A-6F relate to reversal of reprogrammed chromatin, tumorigenicity, and malignant gene expression programs by 6AN.
- [0046] FIG. 6A is a series of graphical representations of data.
- [0047] FIG. 6B is a series of graphical representations of data.
- [0048] FIG. 6C is a series of images of tumor forming assays and related graphical plots.
- [0049] FIG. 6D is a series of images and related graphical plots.
- [0050] FIG. 6E is a series of graphical plots.
- [0051] FIG. 6F is a series of images of tumor forming assays and related graphical plots.
- [0052] FIGS. 7A-7C relate to reprogrammed chromatin across distant metastatic subclones.
- [0053] FIG. 7A is a series of western blot images.
- [0054] FIG. 7B is a series of western blot images.
- [0055] FIG. 7C is a series of western blot images.
- [0056] FIGS. 8A-8E relate to specificity of reprogrammed histone modifications.
- [0057] FIG. 8A is a series of immunohistochemical stains.
- [0058] FIG. 8B is a graphical representation of data.
- [0059] FIG. 8C is a table.
- [0060] FIG. 8D is a series of western blot images.
- [0061] FIG. 8E is a series of western blot images.
- [0062] FIG. 9 is a series of graphical plots relating to enrichment of heterochromatin modifications within LOCKs.
- [0063] FIGS. 10A-10B relate to reprogramming of H3K9Me3 in LOCKs during PDAC subclonal evolution.
- [0064] FIG. 10A is a graphical representation of data.
- [0065] FIG. 10B is a graphical representation of data.
- [0066] FIGS. 11A-11B relate to local reprogramming of DE gene loci within LOCKs.
- [0067] FIG. 11A is a series of graphical representations of data.
- [0068] FIG. 11B is a series of graphical representations of data.
- [0069] FIG. 12 is a series of graphical plots relating to enrichment of euchromatin modifications within ECDs.
- [0070] FIGS. 13A-13E relate to reprogramming of large LOCKs during PDAC evolution.
- [0071] FIG. 13A is a graphical representation of data.
- [0072] FIG. 13B is a graphical representation of data.
- [0073] FIG. 13C is a graphical representation of data.
- [0074] FIG. 13D is a graphical representation of data.
- [0075] FIG. 13E is a graphical representation of data.
- [0076] FIGS. 14A-14F relate to malignant heterogeneity between A38 subclones.
- [0077] FIG. 14A is a graphical representation of data.
- [0078] FIG. 14B is a series of graphical representations of data.
- [0079] FIG. 14C is a series of immunohistochemical stains.
- [0080] FIG. 14D is a series of immunohistochemical stains.
- [0081] FIG. 14E is a series of western blot images.
- [0082] FIG. 14F is a series of images of tumor forming assays and related graphical plots.
- [0083] FIGS. 15A-15C relate to rearrangements targeted to Large LOCKs and ECDs.
- [0084] FIG. 15A is a series of graphical representations of data.
- [0085] FIG. 15B is a series of graphical representations of data.
- [0086] FIG. 15C is a series of graphical representations of data.
- [0087] FIGS. 16A-16B relate to enhanced glucose metabolism with depleted 6PG levels across distant metastases.
- [0088] FIG. 16A is a series of graphical representations of data.
- [0089] FIG. 16B is a series of graphical representations of data.
- [0090] FIGS. 17A-17C relate to 6AN targeting of glucose metabolism and the PGD step of the PPP.
- [0091] FIG. 17A is a series of graphical representations of data.
- [0092] FIG. 17B is a series of graphical representations of data.
- [0093] FIG. 17C is a series of graphical representations of data.
- [0094] FIGS. 18A-18C relate to 6AN selectively modulation of the reprogrammed chromatin state of distant metastatic subclones.
- [0095] FIG. 18A is a series of western blot images.
- [0096] FIG. 18B is a series of western blot images.

[0097] FIG. 18C is a series of western blot images.

[0098] FIGS. 19A-19D relate to 6AN regulated gene expression in LOCK-EI regions.

[0099] FIG. 19A is a series of graphical representations of data.

[0100] FIG. 19B is a series of graphical representations of data.

[0101] FIG. 19C is a series of graphical representations of data.

[0102] FIG. 19D is a series of graphical representations of data.

[0103] FIGS. 20A-20C relate to 6AN selectively blocked tumor formation in distant metastatic subclones.

[0104] FIG. 20A is a series of images of tumor forming assays and related graphical plots.

[0105] FIG. 20B is a series of images of tumor forming assays and related graphical plots.

[0106] FIG. 20C is a series of images of tumor forming assays and related graphical plots.

[0107] FIGS. 21A-21C relates to reprogramming of the TOP2B locus in response to 6AN.

[0108] FIG. 21A is a series of graphical representations of data.

[0109] FIG. 21B is a series of graphical representations of data.

[0110] FIG. 21C is a series of graphical representations of data.

[0111] FIG. 22 is a screen shot illustrating data directly linking loss of large-scale heterochromatic regions as described herein to increased variability of gene expression, allowing for increased phenotypic plasticity. An example is a gene SHC4 that is involved in ERK signaling and tumor invasion and metastasis. Its expression variability statistical index measured by single cell RNA experiments is +1.12 in the A38-5 (epigenomically altered, distant metastatic) line in the paper, and -2.68 in the corresponding A38-41 (epigenomically stable, locally invasive) line, with a FDR p value of 0.00. FIG. 22 represents the data showing the loss of LOCKs over the gene in 38-5.

DETAILED DESCRIPTION OF THE INVENTION

[0112] The present invention is based on the seminal discovery that a prometastatic candidate is epigenomic regulation. The invention is based on discovery of the role of large-scale epigenomic changes during PDAC subclonal evolution and distant metastasis, especially within heterochromatin domains including large organized heterochromatin lysine (K)-9 modified domains (LOCKs) and large DNA hypomethylated blocks. These regions represent selectable targets for large-scale epigenetic reprogramming, since they occupy over half of the genome, partially overlap with one another, and are found in many human cancers including PDAC. The inventors therefore hypothesized that epigenomic dysregulation within these regions could be a major selective force for tumor progression, given the lack of any consistent metastasis-specific driver mutations.

[0113] Before the present systems and methods are described, it is to be understood that this invention is not limited to particular systems, methods, and experimental conditions described, as such systems, methods, and conditions may vary. It is also to be understood that the terminology used herein is for purposes of describing particular

embodiments only, and is not intended to be limiting, since the scope of the present invention will be limited only in the appended claims.

[0114] As used in this specification and the appended claims, the singular forms “a”, “an”, and “the” include plural references unless the context clearly dictates otherwise. Thus, for example, references to “the method” includes one or more methods, and/or steps of the type described herein which will become apparent to those persons skilled in the art upon reading this disclosure and so forth.

[0115] Unless defined otherwise, all 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. Although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of the invention, the preferred methods and materials are now described.

[0116] The present invention provides a method of identifying targets for epigenetic reprogramming comprising detecting large organized heterochromatin lysine (K)-9 modified domains (LOCKs) and large DNA hypomethylated blocks in a sample containing DNA from a subject having cancer. For example, the method applies to a subject that has or is at risk of having PDAC and/or metastasis thereof. In one aspect, the detection comprises analysis of H3K9Me2/3 and/or H4K20Me3.

[0117] As used herein, reprogramming, is intended to refer to a process that alters or reverses the differentiation status of a somatic cell that is either partially or terminally differentiated. Reprogramming of a somatic cell may be a partial or complete reversion of the differentiation status of the somatic cell. In an exemplary aspect, reprogramming is complete wherein a somatic cell is reprogrammed into an iPS cell. However, reprogramming may be partial, such as reversion into any less differentiated state. For example, reverting a terminally differentiated cell into a cell of a less differentiated state, such as a multipotent cell.

[0118] As used herein, pluripotent cells include cells that have the potential to divide in vitro for an extended period of time (greater than one year) and have the unique ability to differentiate into cells derived from all three embryonic germ layers, namely endoderm, mesoderm and ectoderm.

[0119] Somatic cells for use with the present invention may be primary cells or immortalized cells. Such cells may be primary cells (non-immortalized cells), such as those freshly isolated from an animal, or may be derived from a cell line (immortalized cells). In an exemplary aspect, the somatic cells are mammalian cells, such as, for example, human cells or mouse cells. They may be obtained by well-known methods, from different organs, such as, but not limited to skin, brain, lung, pancreas, liver, spleen, stomach, intestine, heart, reproductive organs, bladder, kidney, urethra and other urinary organs, or generally from any organ or tissue containing living somatic cells, or from blood cells. Mammalian somatic cells useful in the present invention include, by way of example, adult stem cells, sertoli cells, endothelial cells, granulosa epithelial cells, neurons, pancreatic islet cells, epidermal cells, epithelial cells, hepatocytes, hair follicle cells, keratinocytes, hematopoietic cells, melanocytes, chondrocytes, lymphocytes (B and T lymphocytes), erythrocytes, macrophages, monocytes, mononuclear cells, fibroblasts, cardiac muscle cells, other known muscle cells, and generally any live somatic cells. In particular embodiments, fibroblasts are used. The term somatic cell, as

used herein, is also intended to include adult stem cells. An adult stem cell is a cell that is capable of giving rise to all cell types of a particular tissue. Exemplary adult stem cells include hematopoietic stem cells, neural stem cells, and mesenchymal stem cells.

[0120] As discussed herein, alterations in methylation patterns occur during differentiation or dedifferentiation of a cell which work to regulate gene expression of critical factors that are ‘turned on’ or ‘turned off’ at various stages of differentiation. As such, one of skill in the art would appreciate that many types of agents are capable of altering the methylation status of one or more nucleic acid sequences of a somatic cell to induce pluripotency that may be suitable for use with the present invention.

[0121] An agent, as used herein, is intended to include any agent capable of altering the methylation status of one or more nucleic acid sequences of a somatic cell. For example, an agent useful in any of the method of the invention may be any type of molecule, for example, a polynucleotide, a peptide, a peptidomimetic, peptoids such as vinylogous peptoids, chemical compounds, such as organic molecules or small organic molecules, or the like. In various aspects, the agent may be a polynucleotide, such as DNA molecule, an antisense oligonucleotide or RNA molecule, such as microRNA, dsRNA, siRNA, stRNA, and shRNA.

[0122] MicroRNA (miRNA) are single-stranded RNA molecules whose expression is known to be regulated by methylation to play a key role in regulation of gene expression during differentiation and dedifferentiation of cells. Thus, an agent may be one that inhibits or induces expression of miRNA or may be a mimic miRNA. As used herein, “mimic” microRNAs which are intended to mean a microRNA exogenously introduced into a cell that have the same or substantially the same function as their endogenous counterpart.

[0123] In various aspects of the present invention, an agent that alters the methylation status of one or more nucleic acid sequences is a nuclear reprogramming factor. Nuclear reprogramming factors may be genes that induce pluripotency and utilized to reprogram differentiated or semi-differentiated cells to a phenotype that is more primitive than that of the initial cell, such as the phenotype of a pluripotent stem cell. Those skilled in the art would understand that such genes and agents are capable of generating a pluripotent stem cell from a somatic cell upon expression of one or more such genes having been integrated into the genome of the somatic cell or upon contact of the somatic cell with the agent or expression product of the gene. As used herein, a gene that induces pluripotency is intended to refer to a gene that is associated with pluripotency and capable of generating a less differentiated cell, such as a pluripotent stem cell from a somatic cell upon integration and expression of the gene. The expression of a pluripotency gene is typically restricted to pluripotent stem cells and is crucial for the functional identity of pluripotent stem cells.

[0124] Several genes have been found to be associated with pluripotency and suitable for use with the present invention as reprogramming factors. Such genes are known in the art and include, by way of example, SOX family genes (SOX1, SOX2, SOX3, SOX15, SOX18), KLF family genes (KLF1, KLF2, KLF4, KLF5), MYC family genes (C-MYC, L-MYC, N-MYC), SALL4, OCT4, NANOG, LIN28, STELLA, NOBOX, POU5F1 or a STAT family gene. STAT family members may include for example STAT1, STAT2,

STAT3, STAT4, STAT5 (STAT5A and STAT5B), and STAT6. While in some instances, use of only one gene to induce pluripotency may be possible, in general, expression of more than one gene is required to induce pluripotency. For example, two, three, four or more genes may be simultaneously integrated into the somatic cell genome as a polycistronic construct to allow simultaneous expression of such genes. In an exemplary aspect, four genes are utilized to induce pluripotency including OCT4, POU5F1, SOX2, KLF4 and C-MYC. Additional genes known as reprogramming factors suitable for use with the present invention are disclosed in U.S. patent application Ser. No. 10/997,146 and U.S. patent application Ser. No. 12/289,873, incorporated herein by reference.

[0125] All of these genes commonly exist in mammals, including human, and thus homologues from any mammals may be used in the present invention, such as genes derived from mammals including, but not limited to mouse, rat, bovine, ovine, horse, and ape. Further, in addition to wild-type gene products, mutant gene products including substitution, insertion, and/or deletion of several (e.g., 1 to 10, 1 to 6, 1 to 4, 1 to 3, and 1 or 2) amino acids and having similar function to that of the wild-type gene products can also be used. Furthermore, the combinations of factors are not limited to the use of wild-type genes or gene products. For example, Myc chimeras or other Myc variants can be used instead of wild-type Myc.

[0126] The present invention is not limited to any particular combination of nuclear reprogramming factors. As discussed herein a nuclear reprogramming factor may comprise one or more gene products. The nuclear reprogramming factor may also comprise a combination of gene products as discussed herein. Each nuclear reprogramming factor may be used alone or in combination with other nuclear reprogramming factors as disclosed herein. Further, nuclear reprogramming factors of the present invention can be identified by screening methods, for example, as discussed in U.S. patent application Ser. No. 10/997,146, incorporated herein by reference. Additionally, the nuclear reprogramming factor of the present invention may contain one or more factors relating to differentiation, development, proliferation or the like and factors having other physiological activities, as well as other gene products which can function as a nuclear reprogramming factor.

[0127] The nuclear reprogramming factor may include a protein or peptide. The protein may be produced from a gene as discussed herein, or alternatively, in the form of a fusion gene product of the protein with another protein, peptide or the like. The protein or peptide may be a fluorescent protein and/or a fusion protein. For example, a fusion protein with green fluorescence protein (GFP) or a fusion gene product with a peptide such as a histidine tag can also be used. Further, by preparing and using a fusion protein with the TAT peptide derived from the virus HIV, intracellular uptake of the nuclear reprogramming factor through cell membranes can be promoted, thereby enabling induction of reprogramming only by adding the fusion protein to a medium thus avoiding complicated operations such as gene transduction. Since preparation methods of such fusion gene products are well known to those skilled in the art, skilled artisans can easily design and prepare an appropriate fusion gene product depending on the purpose.

[0128] In certain embodiments, the agent alters the methylation status of one or more nucleic acid sequences, such as any gene listed in a Table set forth herein.

[0129] Expression profiling of reprogrammed somatic cells to assess their pluripotency characteristics may also be conducted. Expression of individual genes associated with pluripotency may also be examined. Additionally, expression of embryonic stem cell surface markers may be analyzed. As used herein, “expression” refers to the production of a material or substance as well as the level or amount of production of a material or substance. Thus, determining the expression of a specific marker refers to detecting either the relative or absolute amount of the marker that is expressed or simply detecting the presence or absence of the marker. As used herein, “marker” refers to any molecule that can be observed or detected. For example, a marker can include, but is not limited to, a nucleic acid, such as a transcript of a specific gene, a polypeptide product of a gene, a non-gene product polypeptide, a glycoprotein, a carbohydrate, a glycolipid, a lipid, a lipoprotein or a small molecule.

[0130] Detection and analysis of a variety of genes known in the art to be associated with pluripotent stem cells may include analysis of genes such as, but not limited to OCT4, NANOG, SALL4, SSEA-1, SSEA-3, SSEA-4, TRA-1-60, TRA-1-81, or a combination thereof iPS cells may express any number of pluripotent cell markers, including: alkaline phosphatase (AP); ABCG2; stage specific embryonic antigen-1 (SSEA-1); SSEA-3; SSEA-4; TRA-1-60; TRA-1-81; Tra-2-49/6E; ERas/ECATS, E-cadherin; β -III-tubulin; γ -smooth muscle actin (γ -SMA); fibroblast growth factor 4 (Fgf4), Cripto, Dax1; zinc finger protein 296 (Zfp296); N-acetyltransferase-1 (Nat1); ES cell associated transcript 1 (ECAT1); ESG1/DPPA5/ECAT2; ECAT3; ECAT6; ECAT7; ECAT8; ECAT9; ECAT10; ECAT15-1; ECAT15-2; Fth117; Sa114; undifferentiated embryonic cell transcription factor (Utf1); Rex1; p53; G3PDH; telomerase, including TERT; silent X chromosome genes; Dnmt3a; Dnmt3b; TRIM28; F-box containing protein (Fbx15); Nanog/ECAT4; Oct3/4; Sox2; Klf4; c-Myc; Esrrb; TDGF1; GABRB3; Zfp42, FoxD3; GDF3; CYP25A1; developmental pluripotency-associated 2 (DPPA2); T-cell lymphoma breakpoint 1 (Tcl1); DPPA3/Stella; DPPA4; as well as other general markers for pluripotency, for example any genes used during induction to reprogram the cell. iPS cells can also be characterized by the down-regulation of markers characteristic of the differentiated cell from which the iPS cell is induced.

[0131] As used herein, “differentiation” refers to a change that occurs in cells to cause those cells to assume certain specialized functions and to lose the ability to change into certain other specialized functional units. Cells capable of differentiation may be any of totipotent, pluripotent or multipotent cells. Differentiation may be partial or complete with respect to mature adult cells.

[0132] “Differentiated cell” refers to a non-embryonic, non-parthenogenetic or non-pluripotent cell that possesses a particular differentiated, i.e., non-embryonic, state. The three earliest differentiated cell types are endoderm, mesoderm, and ectoderm.

[0133] Pluripotency can also be confirmed by injecting the cells into a suitable animal, e.g., a SCID mouse, and observing the production of differentiated cells and tissues. Still another method of confirming pluripotency is using the subject pluripotent cells to generate chimeric animals and observing the contribution of the introduced cells to different

cell types. Methods for producing chimeric animals are well known in the art and are described in U.S. Pat. No. 6,642, 433, incorporated by reference herein.

[0134] Yet another method of confirming pluripotency is to observe cell differentiation into embryoid bodies and other differentiated cell types when cultured under conditions that favor differentiation (e.g., removal of fibroblast feeder layers).

[0135] In various aspects of the invention, methylation status is converted to an M value. As used herein an M value, can be a log ratio of intensities from total (Cy3) and McrBC-fractionated DNA (Cy5): positive and negative M values are quantitatively associated with methylated and unmethylated sites, respectively.

[0136] In various aspects of the invention large hypomethylated blocks are identified Hypomethylation is present when there is a measurable decrease in methylation. In some embodiments, a DNA block is hypomethylated when less than 50% of the methylation sites analyzed are not methylated. DNA block. Methods for determining methylation states are provided herein and are known in the art. In some embodiments methylation status is converted to an M value. As used herein an M value, can be a log ratio of intensities from total (Cy3) and McrBC-fractionated DNA (Cy5): positive and negative M values are quantitatively associated with methylated and unmethylated sites, respectively. M values are calculated as described in the Examples. In some embodiments, M values which range from -0.5 to 0.5 represent unmethylated sites as defined by the control probes, and values from 0.5 to 1.5 represent baseline levels of methylation.

[0137] Numerous methods for analyzing methylation status of a gene are known in the art and can be used in the methods of the present invention to identify either hypomethylation or hypermethylation of the one or more DMRs. In various embodiments, the determining of methylation status in the methods of the invention is performed by one or more techniques selected from the group consisting of a nucleic acid amplification, polymerase chain reaction (PCR), methylation specific PCR, bisulfite pyrosequencing, single-strand conformation polymorphism (SSCP) analysis, restriction analysis, microarray technology, and proteomics. As illustrated in the Examples herein, analysis of methylation can be performed by bisulfite genomic sequencing. Bisulfite treatment modifies DNA converting unmethylated, but not methylated, cytosines to uracil. Bisulfite treatment can be carried out using the METHYLEASY bisulfite modification kit (Human Genetic Signatures).

[0138] In some embodiments, bisulfite pyrosequencing, which is a sequencing-based analysis of DNA methylation that quantitatively measures multiple, consecutive CpG sites individually with high accuracy and reproducibility, may be used.

[0139] It will be recognized that depending on the site bound by the primer and the direction of extension from a primer, that the primers listed above can be used in different pairs. Furthermore, it will be recognized that additional primers can be identified within the hypomethylated blocks, especially primers that allow analysis of the same methylation sites as those analyzed with primers that correspond to the primers disclosed herein.

[0140] Altered methylation can be identified by identifying a detectable difference in methylation. For example, hypomethylation can be determined by identifying whether

after bisulfite treatment a uracil or a cytosine is present at a particular location. If uracil is present after bisulfite treatment, then the residue is unmethylated. Hypomethylation is present when there is a measurable decrease in methylation.

[0141] In an alternative embodiment, the method for analyzing methylation of a hypomethylated block can include amplification using a primer pair specific for methylated residues within a DMR. In these embodiments, selective hybridization or binding of at least one of the primers is dependent on the methylation state of the target DNA sequence (Herman et al., *Proc. Natl. Acad. Sci. USA*, 93:9821 (1996)). For example, the amplification reaction can be preceded by bisulfite treatment, and the primers can selectively hybridize to target sequences in a manner that is dependent on bisulfite treatment. For example, one primer can selectively bind to a target sequence only when one or more base of the target sequence is altered by bisulfite treatment, thereby being specific for a methylated target sequence.

[0142] Other methods are known in the art for determining methylation status of a hypomethylated block, including, but not limited to, array-based methylation analysis and Southern blot analysis.

[0143] Methods using an amplification reaction, for example methods above for detecting hypomethylation or hypermethylation of one or more hypomethylated blocks, can utilize a real-time detection amplification procedure. For example, the method can utilize molecular beacon technology (Tyagi et al., *Nature Biotechnology*, 14: 303 (1996)) or Taqman™ technology (Holland et al., *Proc. Natl. Acad. Sci. USA*, 88:7276 (1991)).

[0144] Also methyl light (Trinh et al., *Methods* 25(4):456-62 (2001), incorporated herein in its entirety by reference), Methyl Heavy (Epigenomics, Berlin, Germany), or SNUPE (single nucleotide primer extension) (see e.g., Watson et al., *Genet Res.* 75(3):269-74 (2000)) can be used in the methods of the present invention related to identifying altered methylation of DMRs.

[0145] As used herein, the term “selective hybridization” or “selectively hybridize” refers to hybridization under moderately stringent or highly stringent physiological conditions, which can distinguish related nucleotide sequences from unrelated nucleotide sequences.

[0146] As known in the art, in nucleic acid hybridization reactions, the conditions used to achieve a particular level of stringency will vary, depending on the nature of the nucleic acids being hybridized. For example, the length, degree of complementarity, nucleotide sequence composition (for example, relative GC:AT content), and nucleic acid type, for example, whether the oligonucleotide or the target nucleic acid sequence is DNA or RNA, can be considered in selecting hybridization conditions. An additional consideration is whether one of the nucleic acids is immobilized, for example, on a filter. Methods for selecting appropriate stringency conditions can be determined empirically or estimated using various formulas, and are well known in the art (see, e.g., Sambrook et al., *supra*, 1989).

[0147] An example of progressively higher stringency conditions is as follows: 2×SSC/0.1% SDS at about room temperature (hybridization conditions); 0.2×SSC/0.1% SDS at about room temperature (low stringency conditions); 0.2×SSC/0.1% SDS at about 42° C. (moderate stringency conditions); and 0.1×SSC at about 68° C. (high stringency conditions). Washing can be carried out using only one of

these conditions, for example, high stringency conditions, or each of the conditions can be used, for example, for 10 to 15 minutes each, in the order listed above, repeating any or all of the steps listed.

[0148] The degree of methylation in the DNA associated with the DMRs being assessed, may be measured by fluorescent in situ hybridization (FISH) by means of probes which identify and differentiate between genomic DNAs, associated with the DMRs being assessed, which exhibit different degrees of DNA methylation. FISH is described, for example, in de Capoa et al. (*Cytometry*, 31:85-92, 1998) which is incorporated herein by reference. In this case, the biological sample will typically be any which contains sufficient whole cells or nuclei to perform short term culture. Usually, the sample will be a sample that contains 10 to 10,000, or, for example, 100 to 10,000, whole cells.

[0149] Additionally, as mentioned above, methyl light, methyl heavy, and array-based methylation analysis can be performed, by using bisulfate treated DNA that is then PCR-amplified, against microarrays of oligonucleotide target sequences with the various forms corresponding to unmethylated and methylated DNA.

[0150] The term “nucleic acid molecule” is used broadly herein to mean a sequence of deoxyribonucleotides or ribonucleotides that are linked together by a phosphodiester bond. As such, the term “nucleic acid molecule” is meant to include DNA and RNA, which can be single stranded or double stranded, as well as DNA/RNA hybrids. Furthermore, the term “nucleic acid molecule” as used herein includes naturally occurring nucleic acid molecules, which can be isolated from a cell, as well as synthetic molecules, which can be prepared, for example, by methods of chemical synthesis or by enzymatic methods such as by the polymerase chain reaction (PCR), and, in various embodiments, can contain nucleotide analogs or a backbone bond other than a phosphodiester bond.

[0151] The terms “polynucleotide” and “oligonucleotide” also are used herein to refer to nucleic acid molecules. Although no specific distinction from each other or from “nucleic acid molecule” is intended by the use of these terms, the term “polynucleotide” is used generally in reference to a nucleic acid molecule that encodes a polypeptide, or a peptide portion thereof, whereas the term “oligonucleotide” is used generally in reference to a nucleotide sequence useful as a probe, a PCR primer, an antisense molecule, or the like. Of course, it will be recognized that an “oligonucleotide” also can encode a peptide. As such, the different terms are used primarily for convenience of discussion.

[0152] A polynucleotide or oligonucleotide comprising naturally occurring nucleotides and phosphodiester bonds can be chemically synthesized or can be produced using recombinant DNA methods, using an appropriate polynucleotide as a template. In comparison, a polynucleotide comprising nucleotide analogs or covalent bonds other than phosphodiester bonds generally will be chemically synthesized, although an enzyme such as T7 polymerase can incorporate certain types of nucleotide analogs into a polynucleotide and, therefore, can be used to produce such a polynucleotide recombinantly from an appropriate template.

[0153] In another aspect, the present invention includes kits that are useful for carrying out the methods of the present invention. The components contained in the kit depend on a number of factors, including: the particular

analytical technique used to detect methylation or measure the degree of methylation or a change in methylation, and the one or more hypomethylated blocks being assayed for.

[0154] Accordingly, the present invention provides a kit for determining a methylation status of one or more hypomethylated blocks of the invention.

[0155] To examine DNA methylation (DNAm) on a genome-wide scale, comprehensive high-throughput array-based relative methylation (CHARM) analysis, which is a microarray-based method agnostic to preconceptions about DNAm, including location relative to genes and CpG content was carried out. The resulting quantitative measurements of DNAm, denoted with M, are log ratios of intensities from total (Cy3) and McrBC-fractionated DNA (Cy5): positive and negative M values are quantitatively associated with methylated and unmethylated sites, respectively. For each sample, ~4.6 million CpG sites across the genome of iPS cells, parental somatic cells and ES cells were analyzed using a custom-designed NimbleGen HD2 microarray, including all of the classically defined CpG islands as well as all nonrepetitive lower CpG density genomic regions of the genome. 4,500 control probes were included to standardize these M values so that unmethylated regions were associated, on average, with values of 0. CHARM is 100% specific at 90% sensitivity for known methylation marks identified by other methods (for example, in promoters) and includes the approximately half of the genome not identified by conventional region preselection. The CHARM results were also extensively corroborated by quantitative bisulfate pyrosequencing analysis.

[0156] In one aspect of the invention, methylation density is determined for a region of nucleic acid. Density may be used as an indication of a hypomethylated block region of DNA, for example. A density of about 0.2 to 0.7, about 0.3 to 0.7, 0.3 to 0.6 or 0.3 to 0.4, or 0.3, may be indicative of a hypomethylated block (the calculated DNA methylation density is the number of methylated CpGs divided by the total number of CpGs sequenced for each sample). Methods for determining methylation density are well known in the art. For example, a method for determining methylation density of target CpG islands has been established by Luo et al. *Analytical Biochemistry*, Vol. 387:2 2009, pp. 143-149. In the method, DNA microarray was prepared by spotting a set of PCR products amplified from bisulfite-converted sample DNAs. This method not only allows the quantitative analysis of regional methylation density of a set of given genes but also could provide information of methylation density for a large amount of clinical samples as well as use in the methods of the invention regarding iPS cell generation and detection. Other methods are well known in the art (e.g., Holemon et al., *BioTechniques*, 43:5, 2007, pp. 683-693).

[0157] The present invention is described partly in terms of functional components and various processing steps. Such functional components and processing steps may be realized by any number of components, operations and techniques configured to perform the specified functions and achieve the various results. For example, the present invention may employ various biological samples, biomarkers, elements, materials, computers, data sources, storage systems and media, information gathering techniques and processes, data processing criteria, statistical analyses, regression analyses and the like, which may carry out a variety of functions. In addition, although the invention is described in the medical diagnosis context, the present invention may be

practiced in conjunction with any number of applications, environments and data analyses; the systems described herein are merely exemplary applications for the invention.

[0158] Methods for analysis according to various aspects of the present invention may be implemented in any suitable manner, for example using a computer program operating on the computer system. An exemplary analysis system, according to various aspects of the present invention, may be implemented in conjunction with a computer system, for example a conventional computer system comprising a processor and a random access memory, such as a remotely-accessible application server, network server, personal computer or workstation. The computer system also suitably includes additional memory devices or information storage systems, such as a mass storage system and a user interface, for example a conventional monitor, keyboard and tracking device. The computer system may, however, comprise any suitable computer system and associated equipment and may be configured in any suitable manner. In one embodiment, the computer system comprises a stand-alone system. In another embodiment, the computer system is part of a network of computers including a server and a database.

[0159] The software required for receiving, processing, and analyzing biomarker information may be implemented in a single device or implemented in a plurality of devices. The software may be accessible via a network such that storage and processing of information takes place remotely with respect to users. The analysis system according to various aspects of the present invention and its various elements provide functions and operations to facilitate biomarker analysis, such as data gathering, processing, analysis, reporting and/or diagnosis. The present analysis system maintains information relating to methylation and samples and facilitates analysis and/or diagnosis. For example, in the present embodiment, the computer system executes the computer program, which may receive, store, search, analyze, and report information relating to the epigenome. The computer program may comprise multiple modules performing various functions or operations, such as a processing module for processing raw data and generating supplemental data and an analysis module for analyzing raw data and supplemental data to generate a disease status model and/or diagnosis information.

[0160] The procedures performed by the analysis system may comprise any suitable processes to facilitate analysis and/or disease diagnosis. In one embodiment, the analysis system is configured to establish a disease status model and/or determine disease status in a patient. Determining or identifying disease status may comprise generating any useful information regarding the condition of the patient relative to the disease, such as performing a diagnosis, providing information helpful to a diagnosis, assessing the stage or progress of a disease, identifying a condition that may indicate a susceptibility to the disease, identify whether further tests may be recommended, predicting and/or assessing the efficacy of one or more treatment programs, or otherwise assessing the disease status, likelihood of disease, or other health aspect of the patient.

[0161] The analysis system may also provide various additional modules and/or individual functions. For example, the analysis system may also include a reporting function, for example to provide information relating to the processing and analysis functions. The analysis system may

also provide various administrative and management functions, such as controlling access and performing other administrative functions.

[0162] The analysis system suitably generates a disease status model and/or provides a diagnosis for a patient based on raw biomarker data and/or additional subject data relating to the subjects. The data may be acquired from any suitable biological samples.

[0163] The following examples are provided to further illustrate the advantages and features of the present invention but are not intended to limit the scope of the invention. While they are typical of those that might be used, other procedures, methodologies, or techniques known to those skilled in the art may alternatively be used.

Example 1

Large-Scale Epigenomic Reprogramming Links
Anabolic Glucose Metabolism to Distant Metastasis
During the Evolution of Pancreatic Cancer
Progression

Background and Hypothesis

[0164] As discussed previously, one prometastatic candidate is epigenomic regulation. In particular, we wished to investigate the role of large-scale epigenomic changes during PDAC subclonal evolution and distant metastasis, especially within heterochromatin domains including large organized heterochromatin lysine (K)-9 modified domains (LOCKS) and large DNA hypomethylated blocks. These regions could represent selectable targets for large-scale epigenetic reprogramming, since they occupy over half of the genome, partially overlap with one another, and are found in many human cancers including PDAC. The inventors therefore hypothesized that epigenomic dysregulation

within these regions could be a major selective force for tumor progression, given the lack of any consistent metastasis-specific driver mutations.

Results

Reprogramming of Global Epigenetic State During the Evolution of Distant Metastasis

[0165] To test this hypothesis, we first determined whether large-scale changes in epigenetic modifications could be detected during PDAC evolution in patient samples in vivo. We previously collected matched primary and metastatic PDAC lesions from individual patients by rapid autopsy and reported the genetic progression of subclonal evolution by whole exome and sanger sequencing for mutations, paired-end sequencing for rearrangements, and whole genome sequencing in a subset of these samples. These samples represent a unique resource especially suited to study tumor evolution, since they were collected from matched primary and metastatic tumors from the same patient(s), each has been deep sequenced, individual subclones have been identified, and no metastasis-specific driver mutations are present. From these patients, we selected a large panel of diverse PDAC samples to test for global epigenomic reprogramming during subclonal evolution. As summarized in Table 1, these samples were chosen because they represented the diversity of PDAC evolution (different regions of primary tumor paired to peritoneal and distant metastases), each sample represented a sequence-verified (sub)clonal population, patients were both treated and untreated, driver mutations were shared by all subclones in each patient in the absence of metastasis-specific drivers, formalin-fixed tissue was available for immunoassays, frozen tissue was available for whole-genome bisulfate sequencing, and cell lines were available for all other experiments.

TABLE 1

PDAC sample characteristics with LOCK epigenetic changes						
Sample (Clonal Relation ^a)	Disease Present.	Source	Driver Genes (in primary and mets)	Chemo	Assays ^b IHC, Western Blots, ChIP- seq, WGBS	Results ^b LOCK Methylation Change
A124PrF (Founder Clone)	Local- regional	Tissue: Primary Tumor	KRAS CDKN2A SMAD4 ATM	Untreated	K9Me2/3 IHC:	Diffusely Positive
A124PrS (Subclone)	Local- regional	Tissue: Primary Tumor	KRAS CDKN2A SMAD4 ATM	Untreated	K9Me2/3 IHC: WGBS:	Diffusely Positive High (75%, control)
A124Per (Metastasis)	Local- regional	Tissue: Peritoneum	KRAS CDKN2A SMAD4 ATM	Untreated	K9Me2/3 IHC: WGBS:	Diffusely Positive High (74%)
A141PrF (Founder Clone)	Local- regional	Tissue: Primary Tumor	KRAS CDKN2A TP53 MLL3 ARID1B	Untreated	K9Me2/3 IHC:	Diffusely Positive
A141PrS (Subclone)	Local- regional	Tissue: Primary Tumor	KRAS CDKN2A TP53 MLL3 ARID1B	Untreated	K9Me2/3 IHC:	Diffusely Positive

TABLE 1-continued

PDAC sample characteristics with LOCK epigenetic changes						
Sample (Clonal Relation ^a)	Disease Present.	Source	Driver Genes (in primary and mets)	Chemo	Assays ^b IHC, Western Blots, ChIP- seq, WGBS	Results ^b LOCK Methylation Change
A141Per (Metastasis)	Local- regional	Tissue: Peritoneum	KRAS CDKN2A TP53 MLL3 ARID1B	Untreated	K9Me2/3 IHC:	Diffusely Positive
A125PrF (Founder Clone)	Distant metastases	Tissue: Primary Tumor	KRAS CDKN2A TP53 ARID1A	Untreated	K9Me2/3 IHC: WGBS:	Diffusely Positive High (74%)
A125PrS (Subclone)	Distant metastases	Tissue: Primary Tumor	KRAS CDKN2A TP53 ARID1A	Untreated	K9Me2/3 IHC: WGBS:	Positive + Negative Reduced (51%)
A125Lv1 (Metastasis)	Distant metastases	Tissue: Liver	KRAS CDKN2A TP53 ARID1A	Untreated	K9Me2/3 IHC: WGBS:	Diffusely Negative Reduced (57%)
A125Lv2 (Metastasis)	Distant metastases	Tissue: Liver	KRAS CDKN2A TP53 ARID1A	Untreated	K9Me2/3 IHC: WGBS:	Diffusely Negative Reduced (59%)
A132PrF (Founder Clone)	Distant metastases	Tissue: Primary Tumor	KRAS CDKN2A TP53 ATM	Untreated	K9Me2/3 IHC:	Diffusely Positive
A132PrS (Subclone)	Distant metastases	Tissue: Primary Tumor	KRAS CDKN2A TP53 ATM	Untreated	K9Me2/3 IHC:	Positive + Negative
A132Lv (Metastasis)	Distant metastases	Tissue: Liver	KRAS CDKN2A TP53 ATM	Untreated	K9Me2/3 IHC:	Positive + Negative
A38PrF (Founder Clone)	Local- regional + Distant metastases	Tissue: Primary Tumor	KRAS TP53 SMAD4	Gem. Bev.	K9Me2/3 IHC:	Diffusely Positive
A38PrS1 (Peritoneal Precursor Subclone)	Local- regional + Distant metastases	Tissue: Primary Tumor	KRAS TP53 SMAD4	Gem. Bev.	K9Me2/3 IHC:	Diffusely Positive
A38PrS2 (Liver/Lung Precursor Subclone)	Local- regional + Distant metastases	Tissue: Primary Tumor	KRAS TP53 SMAD4 SMARCA2 ^c	Gem. Bev.	K9Me2/3 IHC:	Positive + Negative
A38Lg1 (Metastasis)	Local- regional + Distant metastases	Tissue: Lung	KRAS TP53 SMAD4	Gem. Bev.	K9Me2/3 IHC:	Diffusely Negative
A38Per (Metastasis)	Local- regional + Distant metastases	Cell Line: Peritoneum	KRAS TP53 SMAD4	Gem. Bev.	K9Me2 Western Blot: K9Me2 ChIP-seq: WGBS:	High (100%, cont.) High (100%, cont.) High (79%, cont.)
AsPC1 ^d (N/A)	Local- regional	Cell Line: Ascites	KRAS CDKN2A TP53 SMAD4	Gem.	K9Me2 Western Blot:	High (102%)
HPAFII ^d (N/A) ⁱ	Local- regional	Cell Line: Ascites	KRAS TP53	Gem.	K9Me2 Western Blot:	High (94%)
Capan2 ^d (N/A)	Local- regional	Cell Line: Primary Tumor	KRAS CDKN2A TP53	Gem.	K9Me2 Western Blot:	High (93%)
A2Lg (Metastasis)	Local- regional + Distant metastases	Cell Line: Lung	KRAS TP53	Taxoprex. Gem.	K9Me2 Western Blot:	Reduced (40%)

TABLE 1-continued

PDAC sample characteristics with LOCK epigenetic changes						
Sample (Clonal Relation ^a)	Disease Present.	Source	Driver Genes (in primary and mets)	Chemo	Assays ^b IHC, Western Blots, ChIP- seq, WGBS	Results ^b LOCK Methylation Change
A2Lv (Metastasis)	Local- regional + Distant metastases	Cell Line: Liver	KRAS TP53	Taxoprex. Gem.	K9Me2 Western Blot:	Reduced (50%)
A6Lv (Metastasis)	Local- regional + Distant metastases	Cell Line: Liver	KRAS TP53 MLL3	Gem. Trox.	K9Me2 Western Blot:	Reduced (47%)
A10Lv (Metastasis)	Distant metastases	Cell Line: Liver	KRAS TP53 MLL3	Untreated	K9Me2 Western Blot:	Reduced (61%)
A13Pr1 (Subclone)	Distant metastases	Cell Line: Primary Tumor	KRAS CDKN2A MYC TP53	Untreated	K9Me2 Western Blot: K9Me2 ChIP-seq:	Reduced (64%) Reduced (25%)
A13Pr2 (Subclone)	Distant metastases	Cell Line: Primary Tumor	KRAS CDKN2A MYC TP53	Untreated	K9Me2 Western Blot: K9Me2 ChIP-seq: WGBS:	Reduced (72%) Reduced (51%) Reduced (81%)
A13Lg (Metastasis)	Distant metastases	Cell Line: Lung	KRAS CDKN2A MYC TP53	Untreated	K9Me2/3 Western Blot: K9Me2 ChIP-seq: WGBS:	Reduced (73%) Reduced (53%) Reduced (86%)
A32O (Metastasis)	Local- regional + Distant metastases	Cell Line: Omentum ^e	KRAS TP53	5-FU	K9Me2 Western Blot:	Reduced (71%) Reduced (58%)
A38Lv (Metastasis)	Local- regional + Distant metastases	Cell Line: Liver	KRAS TP53 SMAD4	Gem. Bev.	K9Me2 Western Blot: WGBS:	Reduced (47%) Reduced (67%)
A38Lg (Metastasis)	Local- regional + Distant metastases	Cell Line: Lung	KRAS TP53 SMAD4	Gem. Bev.	K9Me2 Western Blot: K9Me2 ChIP-seq: WGBS:	Reduced (58%) Reduced (48%) Reduced (72%)

Table 1: Summary of H3K9 and DNA methylation changes across tissue and cell line samples. Samples from patients with local-regional spread (peritoneal/ascites) showed relatively high global H3K9/DNA methylation as indicated by multiple assays (right two columns), while samples from patients with distant metastases showed reduced methylation across all assays, which initiated in primary tumors as indicated.

Abbreviations:

Gem. (Gemcitabine), Bev. (Bevacizumab), Taxoprex. (Taxoprexin), Trox. (Troxacitabine).

Superscript Notes

^aClonal origins represent phylogenetic estimates from previously published (ref. 1) and other unpublished (text footnote 1) whole-genome sequencing data.

^bWestern blot data reflect densitometry percentages of H3K9Me2 signals relative to A38Per controls (cont.). Western blots are shown in Supplementary FIG. 1 and the absolute densitometry values are shown in FIG. 1g with p-values included in the figure legend. ChIP-seq data reflect percent of LOCK Mb with reduced H3K9Me2 relative to A38Per controls (cont.), as detailed with Mb and RPKM values in Supplementary Data File 2. WGBS data reflect percent of DNA methylation within LOCKs relative to A124Pr controls (tissues) and A38Per controls (cell lines), as detailed in Supplementary FIGS. 1 and 2.

^cThis metastasis from a chemotherapy-treated patient had a missense mutation in SMARCA2 of unclear significance.

^dThese cell lines were not from the rapid autopsy cohort and rely on previously published genotyping data which may underestimate the driver mutations.

^eThe A32O cell line was isolated from an omental mass lesion in a patient with very aggressive disease including widespread lung metastases, and showed findings similar to the other distant (lung/liver) metastatic subclones.

[0166] We began our analysis with the formalin-fixed tissue samples (totaling 16 uniquely matched, sequence-verified tumor sections, Table 1). We performed immunostains against heterochromatin modifications (e.g., H3K9Me2/3) in order to detect global changes from heterochromatin domains (including LOCKs as defined in Wen et al. (Large histone H3 lysine 9 dimethylated chromatin blocks distinguish differentiated from embryonic stem cells. *Nat Genet* 41, 246-50 (2009)), and McDonald et al. (Genome-scale epigenetic reprogramming during epithelial-to-mesenchymal transition. *Nat Struct Mol Biol* 18, 867-74

(2011))) that might be selectable targets during subclonal evolution. Immunostains revealed diffusely positive (>80%) H3K9Me2/3 staining of PDAC cell nuclei across both primary tumor and peritoneal metastatic subclones from patients who presented with peritoneal carcinomatosis (FIGS. 1A-1B and Table 1). In contrast, samples from patients who presented with distant metastatic disease displayed progressive loss of H3K9Me2/3 during subclonal evolution. This manifested as heterogeneous (mixtures of positive+negative PDAC nuclei) staining in primary tumors followed by either diffusely negative (<20%) staining or

retention of heterogeneous staining in the paired metastases (FIGS. 1C-1D and Table 1). We also observed similar results during subclonal evolution from a patient for which sequence-verified primary tumor subclones that seeded both peritoneal and distant metastases were available. The peritoneal precursor retained diffusely strong staining of heterochromatin modifications as seen in the clone that founded the neoplasm (FIG. 1E, top two panels). In contrast, cell-to-cell heterogeneity of staining patterns emerged in the primary tumor subclone that seeded distant metastasis, followed by diffuse loss of staining at the distant metastatic site (FIG. 1E, bottom two panels). The collective findings across samples from patients with distant metastatic disease thus suggested that reprogramming initiated during subclonal evolution in the primary tumor, and that these changes were inherited or even accentuated in subclones that formed tumors at the distant metastatic sites themselves.

[0167] To expand our analysis to more patient samples and test the generality of our findings, we employed twelve low-passaged cell lines collected from eight patients (Table 1), including a subset that corresponded to the patient tissues above. Cell lines were isolated from nine distant metastatic subclones, a peritoneal metastasis with paired liver and lung metastases that corresponded to the patient presented in FIG. 1E, and two (non-founder) primary tumor subclones matched to a lung metastasis collected from the same patient. Importantly, six of the nine distant metastatic cell lines were previously whole exome sequenced, and mutations present in the cell lines were also present in the corresponding patient tissues as detected by Sanger sequencing. Because rapid autopsy cell lines were largely isolated from patients who presented with distant metastases, additional PDAC samples from other sources of regional disease were also included: malignant ascites fluid from two patients with peritoneal carcinomatosis (AsPC1, HPAFII), and a primary tumor from a long-term survivor without distant metastases (Capan2).

[0168] We then examined whether global changes in chromatin as seen in the patient tissues were maintained in cell lines, which could reflect genome-wide reprogramming events. We began by performing western blots for eight histone modifications with well-understood functions. Comparison of local-regional PDAC samples (A38Per, AsPC1, HPAFII, Capan2) by western blots showed minimal or non-recurrent global changes across all histone modifications tested (FIG. 7A), similar to the evolution of peritoneal carcinomatosis observed in patient tissues. In contrast, distant metastases displayed striking reprogramming of methylation and acetylation that was targeted to specific histone residues (FIG. 7B), including between the paired peritoneal (A38Per) and distant metastatic (A38Lv, A38Lg) subclones that corresponded to the patient presented in FIG. 1E (FIG. 1F). This manifested as recurrent reductions in H3K9Me2/3 and H4K20Me3 that was coupled to hyper-acetylation of H3K9Ac and H3K27Ac in distant metastases (summarized in FIG. 1G and Table 1, shown in FIG. 7B). H3K9 methylation is critical for encoding heterochromatic epigenetic states over large genomic regions including LOCKs, and H3K27Ac encodes gene regulatory elements. Reprogramming appeared specific, as we observed no consistent/recurrent changes in H3K27Me3 or H3K36Me3 across samples (FIG. 1G, FIGS. 7A-7C) and the reprogrammed modifications themselves were not dependent on proliferation rates and could not be induced by PDAC chemotherapy

(FIGS. 8A-8E). Finally, western blots on cell lines isolated from matched primary tumor (A13Pr1, A13Pr2) and distant metastatic (A13Lg) subclones collected from a patient who presented with widespread distant metastases in the absence of regional (peritoneal) spread also showed reductions in H3K9Me3 and H4K20Me3 between the primary tumor subclones, which was retained in the distant metastasis (FIG. 7C), further suggesting that reprogramming initiated in the primary tumor during the evolution of distant metastasis. Thus, in vivo tissue and in vitro cell culture findings across the collective 30 patient samples strongly suggested that global epigenetic state was reprogrammed during the evolution of distant metastasis.

The Epigenomic Landscape of PDAC Subclonal Evolution

[0169] We next wished to map the locations of reprogrammed chromatin modifications across the PDAC genome. To this end, we comprehensively mapped the epigenetic landscape of PDAC evolution with chromatin immunoprecipitation followed by high-throughput sequencing (ChIP-seq) for histone modifications with well-understood functions (heterochromatin: H3K9Me2, H3K9Me3, H3K27Me3; euchromatin: H3K27Ac, H3K36Me3). To capture the diversity of subclonal evolution and malignant progression, ChIP-seq was performed on sequence-verified cell lines isolated from matched subclones, including a peritoneal metastasis (A38Per) matched to a lung metastasis (A38Lg) from the same patient, and two primary tumor subclones (A13Pr1, A13Pr2) that were also matched to a lung metastasis (A13Lg) from the same patient. For each patient, all subclones shared identical driver gene mutations without acquisition of new metastasis-specific drivers (Table 1). We also performed RNA-seq in parallel to identify matched gene expression changes. Finally, we complemented these datasets with whole genome bisulfate sequencing (WGBS) across these cell lines and frozen tumor tissues that corresponded to a subset of the formalin-fixed tumor sections presented in FIGS. 1A-1G. In all, we generated 183 datasets with 19.3×10^9 uniquely aligned sequencing reads, including $>15.0 \times 10^6$ (median: 32.3×10^6) uniquely aligned reads for each ChIP experiment as recommended by ENCODE guidelines (Supplementary Data 1). Experiments were performed as biological replicates, with good correlation between replicates (median correlation coefficient: 0.956; range: 0.746-0.997, Supplementary Data 1). To our knowledge, this represents the first comprehensive genome-wide analysis of epigenetic reprogramming during the evolutionary progression of a human cancer.

[0170] Because global chromatin modifications were stably inherited during the evolution of peritoneal carcinomatosis whereas reprogramming emerged during distant metastasis, we began by comparing the peritoneal subclone against the distant metastatic subclones and their matched primary tumor subclones. This analysis revealed a striking and unexpected degree of genome-wide epigenetic reprogramming that was targeted to thousands of chromatin domains covering $>95\%$ of the PDAC genome. These included heterochromatin regions corresponding to large organized chromatin lysine (K)-modified domains (LOCKs) that occupied approximately half the genome, gene-rich euchromatin domains (ECDs), and a smaller subset of very large LOCK domains that were uniquely reprogrammed compared to other heterochromatin regions. The collective domain characteristics are detailed in Supplementary Data 2,

and comprehensive statistical analyses across experiments are presented in Supplementary Data 3.

[0171] We first analyzed heterochromatin domains defined by strong, broad enrichments of H3K9Me2 and H3K27Me3 with depletion of euchromatin modifications (FIG. 2A, FIG. 9, and Supplementary methods). Thousands of heterochromatin domains were detected in each sample (range: 2,008-3,166) and were organized into large block-like segments (median lengths: 232 Kb-311 Kb) that occupied more than half of the genome in each subclone (average: 61.7% of the genome, range: 54.1-71.7%, Supplementary Data 2). The domain calls were robust as determined by sensitivity analyses across multiple thresholds (Supplementary Data 3), and the called heterochromatin regions themselves overlapped significantly with previously reported LOCK heterochromatin domains (avg: 76.7+/-16.9% overlap; $p < 0.01$ by permutation testing), suggesting that heterochromatin largely corresponded to LOCKs. Similar to immunostain (FIGS. 1A-1D) and western blot data (FIG. 1E-1F), we detected strong H3K9Me2 enrichment across LOCKs in the peritoneal subclone, whereas these same regions displayed global reductions of H3K9Me2 in the distant metastases and their matched primary tumor subclones (FIG. 2A, average: 591 Mb/1,470 Mb; range: 204-1,110 Mb/1,470 Mb; $p < 2.2e-16$ by chi square; Table 1 and Supplementary Data 3). In contrast, high global levels of H3K27Me3 were detected from these regions across all subclones (FIG. 2A), similar to the western blot findings (FIG. 1G). We also detected patient-specific patterns of global H3K9Me3 reprogramming in LOCKs (FIGS. 10A-10B). In patient A38, H3K9Me3 was largely absent from A38Lg LOCKs relative to A38Per (loss of H3K9Me3 from 184/208 Mb, 88.5%, $p < 2.2e-16$). Loss of LOCK-wide H3K9Me3 was also detected between the paired primary tumor subclones in patient A13 (106 Mb/118 Mb, 90.0% in A13Pr2 vs. A13Pr1, $p < 2.2e-16$), and similar to western blot findings (Supplementary FIG. 1C) this change was inherited in the distant metastatic subclone (152 Mb/169 Mb, 90.2% in A13Lg vs. A13Pr1, $p < 2.2e-16$). Finally, localized reprogramming events were also detected specifically within chromatin encoding differentially expressed (DE) genes in LOCKs (FIGS. 11A-11B, Supplementary Data 3). This included reciprocal changes in H3K27Ac and H3K9Me2 over promoters coupled to similar reciprocal changes in H3K36Me3 and H3K27Me3 over gene bodies of LOCK genes that were up- and down-regulated. This suggested that DE genes from LOCKs were situated in specific sub-regions that possessed gene regulatory potential, consistent with hybrid LOCK-euchromatin islands (LOCK-EIs). Collectively, these findings indicated that heterochromatin domains (LOCKs) represented a major target for both global and local chromatin reprogramming events during the evolution of distant metastasis.

[0172] Because LOCKs correspond to a subset of block-like regions that are DNA hypomethylated in pancreatic and other human cancers, we also asked whether DNA methylation changes were targeted to LOCKs during PDAC subclonal evolution. For this analysis, we performed WGBS on all of the cell lines with ChIP-seq data reported above. Because frozen tissues corresponding to the cell lines had been exhausted during previous studies, we selected 7 other frozen tissue samples for in vivo WGBS that were uniquely matched to the same formalin-fixed tissues with IHC data presented in FIG. 1A (A124, local-regional spread) and FIG.

1C (A125, distant metastasis). Normal pancreas was included with frozen tissue samples as an internal control. All samples and results of WGBS with corresponding IHC, western blot, and ChIP-seq findings are summarized in Table 1, and quantified WGBS results with statistical analyses are presented in Tables 2 and 3. These experiments revealed significant reductions in LOCK-wide DNA methylation across cell lines isolated from distant metastases relative to peritoneal carcinomatosis (FIGS. 2B-2C, Table 3). These findings matched the reductions of H3K9Me2 in LOCKs detected by ChIP-seq on the same samples (FIG. 2A), revealing that reprogramming of DNA methylation in hypomethylated block regions is targeted to LOCKs with reprogrammed histone methylation (Table 1). Analysis of the same LOCK regions from the frozen tissue samples also revealed relatively high DNA methylation in LOCKs from patient A124 (peritoneal spread) and the founder clone from patient A125, while the primary tumor and distant metastatic subclone descendants displayed striking loss of DNA methylation that was even more pronounced than that seen in the cell lines (FIGS. 2B-2C, Table 1, and Table 2). We also detected strong, localized DNA hypomethylation from down-regulated DE genes in the hybrid LOCK-EI sub-regions, while up-regulated genes remained hypermethylated with sharp dips at the 5'-ends of genes, similar to H3K9Me2 (FIGS. 11A-11B). Thus, DNA methylation was globally and locally reprogrammed across LOCKs from primary tumor and distant metastatic subclones, similar to histone modifications. Based on the collective immunostain (FIGS. 1A-1E), western blot (FIGS. 1F-1G), ChIP-seq (FIG. 2A), and WGBS (FIGS. 2B-2C) data (Summarized in Table 1), we conclude that a substantial fraction of global reprogramming events was targeted to heterochromatin domains (LOCKs) during the evolution of distant metastasis.

TABLE 2

Percent CpG Methylation levels across LOCK domains detected in frozen tissue samples by WGBS. DNA methylation levels were relatively high in both primary tumor and metastatic tumors from patient A124, who presented with peritoneal carcinomatosis. Similar high levels of DNA methylation were also detected in the founder clone from patient A125, which were significantly reduced in the primary tumor subclone that seeded distant metastases and in the liver metastases themselves. P-values were calculated with paired wilcox tests using a 3% threshold.

Samples	Global LOCK Methylation	Significant Difference
Sample name	% CpG Methylation in	p-value vs. A124Pr
Sample source	LOCKs by WGBS	
A124PrF	75.37%	N/A (Highest Methylation)
Primary Tumor		
A124Per	74.17%	1
Peritoneal Met		
A125PrF	73.65%	1
Primary Tumor #1 (Founder Clone)		
A125PrS	51.12%	2.20E-16
Primary Tumor #2 (Subclone)		
A125Lv1	56.95%	2.20E-16
Liver Met #1		
A125Lv2	59.47%	2.20E-16
Liver Met #2		
Normal Pancreas	67.75%	2.20E-16

TABLE 3

Percent CpG Methylation levels across LOCK domains detected in cell lines by WGBS. DNA methylation levels were highest for the peritoneal subclone A38Per across cell lines. Methylation was significantly reduced in distant metastases from the same patient (A38Lv, A38Lg) and in primary tumor precursors (A13Pr1/2) and the matched lung metastasis from patient A13. P-values were calculated with paired wilcox tests using a 3% threshold.		
Samples Name Source	Global LOCK Methylation % CpG Methylation in LOCKs by WGBS	Significant Difference p-value vs. A38Per
A38Per	78.58%	N/A (Highest Methylation)
Peritoneal Met		
A38Lv	67.39%	2.20E-16
Liver Met		
A38Lg	71.67%	2.20E-16
Lung Met		
A13Pr1	72.46%	2.18E-07
Primary Tumor 1 Subclone		
A13Pr2	73.48%	2.14E-06
Primary Tumor 2 Subclone		
A13Lg	71.30%	2.20E-16
Lung Met		

[0173] We next analyzed reprogramming within ECDs, which were defined by enrichments for global euchromatin modifications H3K27Ac and H3K36Me3 with depletion of heterochromatin modifications (FIG. 12). Similar to heterochromatin, thousands of these domains (range: 1,935-2,318) were partitioned into large, block-like segments (median lengths: 207 Kb-277 Kb) that occupied similar lengths of the genome across subclones (average: 29% of the genome; range: 23.5%-32.0%, Supplementary Data 2). All subclones displayed similar global patterns of modifications within ECDs, including broad H3K36Me3 signals over gene bodies that were flanked by sharp peaks of H3K27Ac and dips in DNA methylation at gene regulatory elements, consistent with actively transcribed euchromatin (FIG. 2D). However, mapping DE genes from RNA-seq data to ECDs (Supplementary Data 3) identified clear patterns of local reprogramming events within chromatin encoding these genes (FIG. 2E, Supplementary Data 3). Genes up-regulated from ECDs acquired increased levels of both H3K36Me3 and H3K27Ac, which could reflect a permissive chromatin state or hyperactive transcription. In contrast, down-regulated genes displayed greatly reduced H3K36Me3 with relatively minor reductions of H3K27Ac, which could reflect an inactive yet poised chromatin state or direct transcriptional repression. Unlike LOCKs, DNA methylation remained stable around DE genes in ECDs (data not shown). Thus, reprogramming in ECDs was largely localized and targeted to H3K27Ac and H3K36Me3 in chromatin encoding DE genes.

[0174] Finally, we also detected patient-specific reprogramming targeted to a unique subset of very large LOCK domains. Although these regions were situated within DNA hypomethylated blocks similar to other LOCKS, they differed in several other respects. First, these domains were substantially larger (median lengths: 730 Kb-1,340 Kb vs. 232-311 Kb for other LOCKs, Supplementary Data 2). Second, they were strongly enriched with H3K9Me3 yet depleted of H3K9Me2/H3K27Me3 (FIGS. 13A-13E and Supplementary Data 2). Third, their abundance was patient-specific: subclones from patient A13 possessed very few of

these domains (range: 50-111 domains covering 1.4-3.5% of the genome) while they occupied a much higher fraction of the A38 genome (range: 226-344 domains covering 14.5-20.6% of the genome, FIG. 13A and Supplementary Data 2). Finally, unlike reprogramming changes detected in other LOCKs (loss of H3K9Me2/3 and DNA methylation), reprogramming in these LOCKs was characterized by loss of H3K9Me3 coupled to increased H3K9Me2 and DNA methylation (FIGS. 13B-13E). Although the functional significance of these findings is uncertain, they could hold implications for patterns of genome instability that emerged during subclonal evolution, as outlined below.

Reprogrammed Chromatin Domains Specify Malignant Heterogeneity

[0175] Subclonal evolution may generate significant phenotypic heterogeneity within an individual patient, and we have hypothesized that such diversity could be encoded by large-scale epigenetic changes similar to those detected above. We therefore wished to investigate in-depth whether reprogrammed chromatin domains might encode heterogeneous malignant properties between PDAC subclones from the same patient. To this end, we selected matched peritoneal and lung metastasis subclones from the same patient (A38Per and A38Lg), performed gene ontology (GO) analyses on reprogrammed LOCK and ECD genes that were differentially expressed between the subclones (Tables 4-7, derived from reprogrammed genes as shown in FIG. 2E and FIGS. 11A-11B), and then tested whether GO results matched actual phenotypic differences measured by experimental assays. This analysis revealed that reprogrammed LOCKs and ECDs encoded substantial phenotypic differences that emerged during subclonal evolution, as described below.

TABLE 4

GO analysis of DE genes that were up-regulated from reprogrammed LOCKs in A38Lg, relative to A38Per. Genes involved in redox balance (oxidation-reduction, NADP) and EMT (cell adhesion, migration) were up-regulated from reprogrammed DE genes in LOCKs (Detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Oxidation-reduction	64	6.0	3.9e-6
Oxidoreductase	56	5.2	4.7e-6
EGF-like domain	28	2.6	6.4e-5
Transferase	105	9.8	1.0e-4
NADP	21	2.0	1.7e-4
Cell Adhesion	41	3.8	1.8e-4
Cell Migration	31	2.9	2.7e-4
Cell Morphogenesis	29	2.7	3.8e-4
Mitochondrion	67	6.3	4.0e-4
Acetylation	174	16.3	5.0e-4

TABLE 5

GO analysis of DE genes that were down-regulated from reprogrammed LOCKs in A38Lg, relative to A38Per. Genes involved in differentiation state (cell adhesion, development, epithelial genes), immune regulation (immune response, cytokines, inflammation), and response to environmental cues (transmembrane signaling, extracellular matrix, secretion, locomotion) were down-regulated from reprogrammed LOCKs (Detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Signal	399	32.4	1.5e-31
Glycoprotein	488	40.0	1.0e-30
Disulfide Bond	352	28.5	1.0e-25
Secreted	217	17.6	1.5e-18
Membrane	582	47.2	1.7e-15
Polymorphism	951	77.1	1.5e-13
Immune Response	106	8.6	1.6e-13
Immunoglobulin Domain	75	6.1	8.5e-11
Cytokine-Cytokine Receptor Interaction	54	4.4	2.3e-10
Ion Channel	55	4.5	2.1e-9
Inflammatory Response	55	4.5	7.6e-9
Cell Adhesion	93	7.5	1.4e-8
Developmental Protein	98	7.9	4.4e-8
Cell Motion	69	5.6	4.7e-8
Transmembrane Protein	82	6.7	3.4e-7
Protease Inhibitor	25	2.0	3.9e-7
Response to Wounding	71	5.8	7.0e-7
Extracellular Matrix	40	3.2	1.2e-6
Epithelial Cell Differentiation	28	2.3	1.6e-6

TABLE 6

GO analysis of DE genes that were up-regulated from reprogrammed ECDs in A38Lg, relative to A38Per. Genes involved in post-translational modifications, cell cycle control, DNA repair, response to stress, and DNA/RNA/protein biosynthesis were up-regulated from reprogrammed ECDs (Detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Acetylation	622	21.7	1.1e-59
Phosphoprotein	1290	45.0	5.1e-53
Cell Cycle	154	5.4	3.4e-30
Mitotic Cell Cycle	137	4.8	4.9e-30
Organelle Fission	95	3.3	3.7e-25
DNA Metabolic Process	157	5.5	9.4e-25
DNA Repair	95	3.3	1.4e-17
Response to DNA Damage Stimulus	113	3.9	3.8e-17
DNA Replication	68	2.4	2.7e-14
Cellular Response to Stress	144	5.0	3.1e-14
Protein Biosynthesis	62	2.1	2.8e-12
ATP Binding	256	8.9	1.3e-11
Ribonucleoprotein	78	2.7	4.1e-11
Nucleotide Binding	309	10.7	4.3e-11
Translation	86	3.0	2.5e-9
ncRNA Metabolic Process	66	2.3	3.5e-9
Mitochondrion	167	5.8	4.3e-9
DNA Recombination	39	1.4	4.6e-9
Microtubule-based Process	70	2.4	5.9e-9

TABLE 7

GO analysis of DE genes that were down-regulated from reprogrammed ECDs in A38Lg, relative to A38Per. Genes involved in oncogenic signal transduction cascades (Sh3 domains, transmembrane proteins, kinases, Ras signaling), cell motion (wounding, migration, locomotion), and cell death control (apoptosis) were down-regulated from reprogrammed ECDs (Detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Alternative Splicing	1195	49.0	1.5e-16
Sh3 Domain	65	2.7	6.8e-10
Phosphoprotein	1112	45.0	1.3e-9
Membrane	964	39.4	1.4e-9
Response to Wounding	117	4.8	9.0e-8
Pleckstrin Homology	71	2.9	2.8e-7
Cell Migration	49	2.0	3.4e-7
Small GTPase Signal Transduction	65	2.7	3.9e-7
Locomotion	52	2.1	1.4e-6
Intracellular Signaling Cascade	227	9.3	3.5e-6
Tyrosine Protein Kinase	34	1.4	5.4e-6
Regulation of Apoptosis	153	6.2	1.1e-5
Protein Kinase Cascade	81	3.3	1.4e-5
Protein Amino Acid Phosphorylation	130	5.3	1.8e-5
Ankyrin Repeat	56	2.3	3.7e-5
Ras Protein Signal Transduction	51	2.1	4.6e-5

[0176] First, a large number of DE genes involved in redox (oxidation-reduction) balance were up-regulated from reprogrammed LOCKs in A38Lg (Table 4, Supplementary Data 3). This subclone was accordingly highly resistant to H₂O₂-mediated oxidative stress (FIG. 3A) and possessed higher oxidoreductase activity and NADPH levels than A38Per (FIGS. 14A-14B). Second, genes encoding differentiation state (epithelial vs. EMT) were reciprocally expressed from A38Per and A38Lg LOCKs (Table 5, Supplementary Data 3), and we confirmed several well-known epithelial and EMT expression changes (e.g., CDH1/E-cadherin, CDH2/N-cadherin) at the protein level by western blots (FIG. 3B). Further consistent with GO results, A38Per maintained well-differentiated (epithelial) morphology while A38Lg was poorly differentiated (EMT-like) across multiple in vitro culture conditions (FIG. 14C), and immunofluorescence experiments showed that EMT emerged in the primary tumor subclone that seeded the A38Lg metastasis in vivo (FIG. 14D). We also note that immune-related genes were differentially expressed from reprogrammed LOCKs (Table 5), which could hold implications for PDAC immunotherapy. Third, genes involved in DNA repair and cell stress responses were significantly up-regulated in ECDs from A38Lg, including genes crucial for maintenance of genome integrity (Fanconi anemia complex, non-homologous end joining, and the TOP2B/OGG1/KDM1A complex, among others, Table 6, Supplementary Data 3). This subclone was accordingly highly resistant to PDAC chemotherapy (gemcitabine) compared to A38Per (FIG. 3C), and western blots showed hyper-phosphorylation of histone H2AX S139 (γH2AX, a signature of activated DNA repair pathways, FIG. 3D). Fourth, genes involved in oncogenic signal transduction cascades were down-regulated in ECDs from A38Lg, especially KRAS/ERK-related genes (Table 7, Supplementary Data 3). Indeed, A38Lg showed loss of phosphorylated ERK (FIG. 3E), resistance to ERK inhibition (FIG. 3F), and minimal response to knock-down of oncogenic KRAS in 3D tumor forming assays (FIG. 3G, FIGS. 14E-14F), despite possessing identical

KRAS^{G12V} mutations as A38Per. Finally, mapping previously reported rearrangements from this patient to chromatin domains revealed that rearrangements were preferentially targeted to ECDs and the small subset of uniquely reprogrammed large LOCK domains, whereas other LOCKs were strongly depleted (FIGS. 15A-15C).

[0177] Thus, reprogrammed chromatin domains collectively specified malignant gene expression programs, divergent phenotypic properties, and patterns of genome instability that emerged during subclonal evolution in patient A38. This patient was unusual in having received chemotherapy prior to tissue harvesting and had a missense mutation in SMARCA2 of unclear significance (CID, unpublished observations), and thus in this case epigenetic selection may have occurred downstream of a genetic driver. Although the nature and extent of such findings will certainly vary among patients, they imply that PDAC is capable of acquiring substantial epigenetic and malignant diversity during subclonal evolution, even in the same cancer from the same patient.

Anabolic Glucose Metabolism Controls Epigenetic State and Tumorigenicity

[0178] We next asked whether a recurrent, metastasis-intrinsic pathway might have been selected for during subclonal evolution to exert upstream control over global epigenetic state and tumorigenic potential. Several recent studies have linked nutrient status and metabolic activity to global levels of histone modifications. Because distant metastases in the rapid autopsy cohort were largely isolated from organs (liver, lung) that provide a rich supply of glucose, we asked whether reprogrammed chromatin and tumorigenicity in these subclones might have evolved a dependence on specific aspects of glucose metabolism.

[0179] Altered glucose metabolism (i.e., Warburg effect) is a well-known property of neoplastic and highly proliferative cells. Although most of our metastatic subclones actually displayed modest proliferative rates in culture (e.g., FIGS. 8A-8E) and in vivo, we nonetheless asked whether distant metastases might have acquired further adaptations in glucose metabolism. Surprisingly, relative to proliferative (immortalized) normal HPDE cells and local-regional PDAC samples, glucose strongly stimulated metabolic (oxidoreductase) activity across distant metastatic subclones (FIG. 4A), and glucose was accordingly required for these subclones to withstand oxidative stress (FIGS. 4B-4C). Distant metastases also hyper-consumed glucose, as we detected elevated glucose uptake and lactate secretion in distant metastases and their precursors relative to peritoneal carcinomatosis (FIG. 16A). To determine if excess glucose uptake was specifically incorporated into downstream metabolic pathways, we selected paired peritoneal and distant metastatic subclones from the same patient, incubated them with ¹³C[1-2]-labeled glucose, and measured glucose incorporation into metabolic products with liquid chromatography followed by high resolution mass spectrometry (LC-HRMS). These experiments revealed elevated incorporation of both C1- and C1,2-labeled glucose into lactate and nucleotides in the distant metastasis (FIGS. 4D-4E), consistent with enhanced glucose entry into both glycolysis and the pentose phosphate pathway (PPP).

[0180] We next asked whether distant metastases might have evolved a dependence on specific enzymatic steps in either of these glucose-driven pathways, which we hypoth-

esized would manifest as severe depletion of metabolite substrate secondary to hyper-consumption. To test this, we surveyed glycolytic and PPP metabolite profiles across a diverse panel of samples including HPDE cells, peritoneal carcinomatosis, distant metastases, and primary tumor precursor subclones. Analysis of all detected glycolytic and pentose phosphate metabolites (FIG. 16B) revealed a striking, recurrent depletion of 6-phosphogluconic acid (6PG) across distant metastases and their precursors (FIG. 4F). 6PG is the substrate for 6-phosphogluconate dehydrogenase (PGD), an enzyme involved in anabolic glucose metabolism that operates within the oxidative branch of the PPP.

[0181] Glucose may enter the PPP via the oxidative (oxPPP) or the non-oxidative (noxPPP) branch of the pathway, which are thought to be uncoupled. Although some studies in other cancers have suggested that PGD is an important oncogene, it is KRAS-mediated noxPPP activation that drives primary tumor growth in mouse models of PDAC. Because KRAS and other driver mutations are acquired early in PDAC progression and shared by all subclones that evolve thereafter, we hypothesized that PGD dependence might have been selected for specifically during the evolution of distant metastasis to maintain reprogrammed chromatin and tumorigenicity. Glucose deprivation, RNAi against PGD, and 6-aminonicotinamide (6AN, a nicotinamide antimetabolite prodrug reported to preferentially inhibit PGD) had no effect on global chromatin modifications in the peritoneal subclone, while all treatments reversed the reprogrammed chromatin state of the paired lung metastasis from the same patient (FIG. 5A). PGD loss-of-function appeared specific, as PGD knockdown did not alter expression of KRAS or other PPP components (FIG. 5B).

[0182] We next asked whether PGD knockdown might affect intrinsic tumor forming capacity across a larger panel of subclones. Despite their aggressive behavior in patients, distant metastatic subclones were unable to effectively form metastatic tumors in immunodeficient mice, and PGD RNAi was not toxic to any subclones grown in routine 2-D cultures (data not shown). To bypass these limitations, we treated cells with RNAi and used 3-D matrigel tumor-forming assays to measure the effects of PGD knockdown on intrinsic tumor-forming capacity. PGD RNAi had minimal effect on the ability of HPDE cells to form spheres or local-regional PDACs to form tumors by these assays (FIG. 5C). Remarkably, PGD RNAi universally interfered with the ability of distant metastatic subclones to form tumors (FIG. 5D). These findings suggested that PGD might represent a therapeutic target with selectivity for PDAC distant metastasis. Because 6AN could represent a lead compound for future design of PGD targeted therapies, we stringently tested it for activity against distant metastases with metabolomics, western blots, multiple 3D tumorigenic assays, RNA-seq, and ChIP experiments.

[0183] 6AN treatments slowed rates of glucose consumption and lactate secretion with no effect on glutamine consumption or glutamate secretion in distant metastatic and precursor subclones (FIG. 17A), and 6AN reversed the previously detected high incorporation of glucose into lactate and nucleotides (FIG. 17B). Furthermore, steady state levels of glucose and metabolites directly upstream of the PGD reaction were dramatically elevated in response to 6AN with corresponding reductions in downstream metabo-

lites (FIG. 17C), which is consistent with strong PGD inhibition as previously reported by others.

[0184] We next tested the effects of 6AN on epigenetic state. Strikingly, 6AN treatments quantitatively reversed several reprogrammed chromatin modifications across distant metastatic subclones with minimal effect on normal cells or local-regional PDACs (FIGS. 18A-18B; summarized in FIGS. 6A-6B), and this effect persisted upon removal of 6AN from the media (FIG. 18C). Because these changes mirrored aspects of LOCK reprogramming, we examined the chromatin state of LOCK DE genes regulated by 6AN, as identified by RNA-seq (Supplementary Data 3, 4). This revealed that DE genes were located within the reprogrammed hybrid LOCK-EI regions that possessed strong H3K27Ac and H3K36Me3, low H3K27Me3, and sharp 5'-depletion (dips) of H3K9Me2 (FIG. 19A, Supplementary Data 3). ChIP-seq experiments on control and 6AN-treated A38Lg cells further showed that the quantitative increase of global H3K9Me2 was targeted to LOCK regions that were reprogrammed in A38Lg vs. A38Per (FIG. 19B), while the reduced H3K27Ac was specifically targeted to genes repressed from LOCKs with no effect on other LOCK genes or ECD-regulated genes (FIG. 19C). Levels of H3K27Me3 remained stable across all regions in response to 6AN (FIG. 19B, FIG. 19D), similar to western blot findings. Collectively, these experiments demonstrated that 6AN selectively and quantitatively targeted several chromatin changes within LOCKs that emerged during the evolution of distant metastasis.

[0185] Because 6AN modulated the global epigenetic state, we hypothesized that it might also selectively block tumorigenic potential in distant metastatic subclones, similar to PGD knockdown experiments. Strikingly, 6AN selectively and strongly blocked tumor formation in distant metastatic and primary tumor precursor subclones but not local-regional PDACs across multiple 3D tumorigenic experimental platforms, including suspension tumorsphere assays (Supplementary FIGS. 20A-20C), matrigel tumor forming assays (FIG. 6B), and injection of PDAC cells into organotypic stroma that recapitulates aspects of in vivo patient tumors (FIG. 6C). Thus, like PGD knockdown, chemical inhibition of PGD by 6AN selectively blocked the tumorigenic potential of distant metastatic subclones.

[0186] We next examined our RNA-seq datasets to explore whether the above findings might be linked to regulation of malignant gene expression programs. Remarkably, over half (952/1832, 52%, Supplementary Data 4) of 6AN down-regulated genes from A38Lg corresponded to genes that were over-expressed in this subclone (compared to the peritoneal subclone from the same patient). In addition, a large fraction of 6AN up-regulated genes also matched DE genes that were repressed (914/2122, 42%, Supplementary Data 4). Even more striking, nearly one-third (255/891, 29%, Supplementary Data 4) of recurrently over-expressed genes across distant metastatic subclones were down-regulated by 6AN. Comparative GO analyses on these gene subsets produced overlapping results that were strongly enriched for cancer-related functions, including mitotic cell cycle control, acetylation, chromosome stability, DNA repair, cell stress responses, and anabolic/biosynthetic activities (Tables 8-10).

TABLE 8

GO Analysis of Genes that were both Recurrently Over-Expressed in Distant Metastases and Down-Regulated by 6AN, Detected by RNA-seq (Detailed in Supplementary Data 3).			
Go Terms	# of Genes	% of Genes	P-value
Nucleus	112	40.0	1.4e-19
Phosphoprotein	150	53.6	1.3e-18
Acetylation	80	28.6	2.7e-16
DNA Metabolic Process	36	12.9	1.2e-15
Cell Cycle	39	13.9	3.4e-12
M-phase	29	8.9	1.8e-11
DNA Repair	22	7.9	2.8e-10
DNA Replication	18	6.4	8.5e-10
Cellular Response to Stress	27	9.6	5.0e-8
Ribonucleotide Biogenesis	13	4.6	6.3e-6

TABLE 9

GO analysis of recurrently over-expressed genes detected by RNA-seq in distant metastatic subclones and primary tumor precursors, relative to peritoneal carcinomatosis (detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Acetylation	217	20.7	8.0e-28
Phosphoprotein	428	40.0	5.3e-24
Protein Biosynthesis	36	3.4	1.4e-14
Nucleus	256	24.4	1.8e-13
Ribonucleoprotein	41	3.9	1.4e-12
Translation	47	4.5	1.5e-12
DNA Metabolic Process	53	5.1	4.5e-9
Cell Cycle	65	6.2	3.4e-7
Mitochondria	64	6.1	5.4e-7
DNA Repair	31	3.0	5.2e-6
Nitrogen Compound Biosynthesis	33	3.1	1.1e-5
Nucleotide Binding	102	9.7	1.4e-6
M-phase	33	3.1	1.4e-5
Cellular Response to Stress	47	4.5	2.5e-5
Transit Peptide	39	3.7	3.3e-5
Nucleotide Biosynthetic Process	22	2.1	5.1e-5
ATP Binding	82	7.8	5.3e-5
DNA Replication	22	2.1	7.0e-5
WD40 Repeat	24	2.3	1.4e-4

TABLE 10

GO analysis of DE genes detected by RNA-seq that were down-regulated in response to 6AN, compared to DMSO control cells (A38Lg subclone, detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Cell Cycle	226	12.2	2.5e-57
Acetylation	465	25.2	5.6e-57
Phosphoprotein	935	50.6	2.5e-55
M-phase	135	7.3	1.2e-52
DNA Metabolic Process	153	13.2	2.6e-40
Nucleus	582	31.5	1.4e-34
DNA Replication	82	4.4	1.0e-33
Chromosome Segregation	45	2.4	3.1e-24
DNA Repair	88	4.5	5.6e-24
Cytoplasm	444	24.0	3.5e-23
ATP Binding	221	12.0	4.1e-22
Cellular Response to Stress	125	6.8	1.6e-19
Chromosome Organization	107	5.8	9.3e-17
Microtubule-based Process	70	3.8	3.6e-16
Nucleotide Binding	244	13.2	4.2e-16
Cytoskeleton	113	6.1	3.2e-15

TABLE 10-continued

GO analysis of DE genes detected by RNA-seq that were down-regulated in response to 6AN, compared to DMSO control cells (A38Lg subclone, detailed in Supplementary Data 3).			
GO Terms	# of Genes	% of Genes	P-value
Ubl Conjugation	103	5.6	9.5e-12
DNA Recombination	36	1.9	1.6e-11
Macromolecular Complex Assembly	115	6.2	3.0e-10

[0187] The above findings led us to hypothesize that 6AN-ablation of tumorigenicity in distant metastatic subclones might be mediated through epigenetic control of cancer-related genes important to maintain tumorigenic capacity. To validate this, we selected two candidate genes for in-depth experiments: N-cadherin (CDH2) and topoisomerase 213 (TOP2B). CDH2 and TOP2B are both thought to be important for cancer progression, are not known to be mutated in PDAC, can be therapeutically targeted, were recurrently over-expressed across distant metastatic and primary tumor precursor subclones by RNA-seq (Supplementary Data 3), and were selectively repressed by 6AN which we confirmed with RT-PCR (FIG. 6E top panels). Furthermore, CDH2 was located within a reprogrammed LOCK targeted by 6AN, and TOP2B was located immediately adjacent to a LOCK boundary. ChIP-qPCR assays performed on control and 6AN treated cells showed nearly identical enrichments for H3K9Me2 and H3K27Ac across these gene loci in the peritoneal subclone (FIG. 6E left panels and FIG. 21A). In contrast, 6AN treatments on the matched lung metastasis from the same patient resulted in enrichment of H3K9Me2 across both loci with concordant reductions of H3K27Ac over the CDH2 genic region (FIG. 6E right panels, FIG. 21A). This strongly suggested that a major downstream effect of 6AN treatments was epigenetic repression of over-expressed cancer genes. We therefore performed RNAi experiments to test whether knockdown of these genes might be important to selectively maintain tumorigenicity. Indeed, RNAi selectively blocked 3D tumor formation in distant metastatic and precursor subclones that over-expressed CDH2 and TOP2B, with no effect on HPDE cells or peritoneal carcinomatosis (FIG. 6F). Collectively, these targeted validation studies strongly supported conclusions inferred from the sequencing data, in that inhibition of PGD/oxPPP by 6AN selectively targeted gene expression, epigenetic state, and downstream tumorigenic functions of over-expressed cancer genes (CDH2/TOP2B).

[0188] As detailed in FIGS. 1A-1G, a global epigenetic reprogramming occurred during the evolution of distant metastasis. The immunohistochemical (IHC) stains against H3K9Me2/3 performed on tumor sections from 6 subclones collected from two patients who presented with widespread peritoneal carcinomatosis (a: patient A124, b: patient A141) showed similar strong nuclear staining across all primary tumor and peritoneal subclones. Similar stains on 6 subclones from two patients who presented with widespread distant metastases (c: patient A125, d: patient A132) showed progressive loss of nuclear staining that initiated in primary tumor subclones that seeded metastases (middle panel) and was further lost (c) or stably inherited (d) in the liver metastases. Scale bars=100 μ m for IHC, 20 μ m for IF. IHC against the indicated modifications performed on tumor sections representing 4 paired subclones from a patient

(patient A38) that presented with both peritoneal carcinomatosis and distant metastases shows that the peritoneal precursor subclone in the primary tumor that seeded carcinomatosis inherited strong nuclear staining of heterochromatin modifications as seen in the parental clone that founded the neoplasm. In contrast, the primary tumor precursor subclone that seeded distant metastases showed cell-to-cell variation in staining, with complete loss of staining in the paired lung metastasis. Staining for the euchromatin modification H3K36Me3 remained stable across all subclones. f, Similar to IHC on tissues (e), western blots on cell lines collected from the peritoneal subclone (Per), liver metastasis, and lung metastasis from patient A38 also showed loss of heterochromatin modifications in distant metastatic subclones, with corresponding increased acetylation. Levels of H3K27Me3 and H3K36Me3 did not differ between subclones. g, Densitometry summary of western blot findings for the indicated histone modifications across cell lines from distant metastatic subclones compared to peritoneal carcinomatosis (n=8 biological replicates, error bars=s.e.m., *p<0.01).

[0189] As detailed in FIGS. 2A-2E, an epigenomic reprogramming of chromatin domains during PDAC subclonal evolution was observed. Representative (left panels) and total summarized (right panels) ChIP-seq experiments revealed loss of H3K9Me2 from LOCKs between peritoneal (A38Per) and distant metastatic and primary tumor precursor subclones (others). H3K27Me3 remained strong in all subclones. Bisulfite-seq data on cell lines (A38, A13, top panel) and frozen tissue samples (A124, A125 panels) showed that samples from local regional spread and parental clones (A38Per, A124PrF, A124Per, A125PrF) possessed hypermethylated LOCKs. In contrast, distant metastatic subclones (A125Lv1/2, A13Lg, A38Lv, A38Lg) and their primary tumor subclones (A125PrS, A13Pr1, A13Pr2) showed hypomethylation of DNA across the same LOCK regions. Global levels of H3K36Me3, H3K27Ac, and DNA methylation within ECDs did not show any clear differences between subclones. In contrast, distant metastatic subclones and primary tumor subclones displayed local reprogramming of H3K36Me3 and H3K27Ac specifically over DE genes within ECDs, compared to the same DE genic ECD regions from A38Per.

[0190] As detailed in FIGS. 3A-3G, reprogrammed chromatin domains encode divergent malignant properties. A38Lg was remarkably resistant to H₂O₂ treatments compared to A38Per. MTT signals reflect cell viability normalized to untreated controls. n=4 technical replicates, *p<0.03. Western blots for proteins involved in epithelial and EMT differentiation were differentially expressed between A38Per and A38Lg, as predicted by GO analyses of reprogrammed DE genes from LOCKs. A38Lg was completely resistant to gemcitabine compared to A38Per, as predicted by GO analyses of reprogrammed DE genes from ECDs. MTT signals reflect cell viability normalized to untreated controls. n=4 technical replicates, *p<0.01. A38Lg possessed elevated levels of γ H2AX by western blot, consistent with activation of DNA repair pathways. Western blots showed that A38Lg lost hyper-phosphorylated ERK and f, was resistant to ERK targeted therapy, compared to A38Per. MTT signals reflect cell viability, normalized to untreated controls. n=4 technical replicates, *p<0.03. A38Lg also lost

sensitivity to KRAS knockdown by Matrigel®3D tumor forming assays, compared to A38Per. n=4 technical replicates, *p<0.01.

[0191] As detailed in FIGS. 4A-4F, hyperactive glucose metabolism and 6PG depletion were observed in distant metastatic subclones. MTT assays performed on equal numbers (20K) of viable, growth-arrested cells from the indicated subclones showed greatly elevated signal (oxidoreductase activity) across distant metastatic subclones, compared to HPDE and local-regional PDAC samples (n=4 technical replicates for each, error bars=s.d.m., *p<10⁻⁵). Normalized cell counts for the indicated samples incubated with (+) or without (-) 10 mM glucose and treated with 1 mM H₂O₂ as indicated (+, -) for 24h showed that normal HPDE cells were sensitive to H₂O₂ under either glucose condition (as expected), whereas local-regional PDAC samples were resistant to H₂O₂ irrespective of glucose availability (n=3 technical replicates for each, error bars=s.d.m.). In contrast, distant metastatic subclones were sensitive to H₂O₂ when glucose was not present in the media (n=3 technical replicates for each, error bars=s.d.m., *p<0.001). Simplified schematic of ¹³C-(1,2)-labeled glucose flow through glycolysis and the PPP. Glucose that enters the oxidative branch of the PPP has one labeled carbon cleaved during conversion of 6PG to Ru5P (m+1), whereas glucose that travels through glycolysis or the non-oxidative PPP retains both labeled carbons (m+2). Note that cross-talk allows glucose with either labeling pattern to re-enter the other pathway and incorporate. LC-MS for nucleotides and lactate showed that these downstream metabolites acquired greatly elevated ¹³C-1,2 labels from glucose in the lung metastasis from patient A38 (A38Lg), compared to its paired peritoneal subclone (A38Per, n=3 biological replicates, error bars=s.d.m., *p<0.01). Steady state LC-HRMS measurements for 6PG showed either complete (ND: not detected) or near complete loss of metabolite across distant metastases and their precursors compared to peritoneal carcinomatosis and HPDE cells.

[0192] As detailed in FIGS. 5A-5D, a PGD-dependence in distant metastatic subclones was observed. Western blots against indicated histone modifications performed on paired peritoneal (A38Per) and distant metastatic (A38Lg) subclones from the same patient showed that global levels of reprogrammed H3K9Me2/3 and acetylation in A38Lg were reversed by removal of glucose from the media (left panel), PGD RNAi (middle panel), and 6AN treatments (right panel). Western blots on A38Lg indicated that PGD knockdown by RNAi did not perturb expression of other PPP components or KRAS. PGD RNAi did not affect the ability of normal HPDE cells or local-regional PDAC samples to form tumors in 3D Matrigel® assays (representative photomicrographs shown with quantified numbers of tumors/well, n=4 technical replicates for each, error bars=s.d.m.). In contrast, PGD RNAi significantly reduced tumor formation across all distant metastatic subclones that were available for testing from the rapid autopsy cohort (n=4 technical replicates for each, error bars=s.d.m., *p<0.01). Scale bars: 200 μm.

[0193] As illustrated in FIGS. 6A-6F, a reversal of reprogrammed chromatin, tumorigenicity, and malignant gene expression programs by 6AN was observed. Densitometry summary of western blots performed on 8 biological replicates, error bars=s.e.m., *p<0.01. 6AN selectively reversed reprogrammed H3K9Me2 and acetylation across most dis-

tant metastatic subclones, with minimal or non-recurrent effects on H3K27Me3 or H4K20Me3. Densitometry summary of western blots performed on 6 biological replicates, error bars=s.e.m. 6AN had minimal effects on histone modifications across normal (HPDE, fibroblast) or local-regional PDAC samples. 6AN ablated tumor formation in 3D Matrigel® assays (n=4 technical replicates for each, error bars=s.d.m., *p<0.01, scale bars: 200 μm) and 3D tumorsphere assays across distant metastatic subclones. 6AN had minimal effects on local-regional PDAC samples by either assay. 6AN also blocked the ability of distant metastatic subclones to form tumors when injected into 3D organotypic stromal cultures (n=3 technical replicates for each, error bars=s.d.m., *p<0.05; scale bars: 200 μm). Real-time RT-PCR (top panels) showed that the distant metastatic subclone (A38Lg) over-expressed CDH2 relative to the peritoneal subclone (A38Per) from the same patient, and that expression was repressed by 6AN (n=4 technical PCR replicates from two biological replicate experiments, error bars: s.d.m., *p=0.002). ChIP assays for H3K9Me2 and H3K27Ac with PCR primers (location indicated by numbers) spaced across the 1.4 Mb chromatin domain showed that 6AN induced spreading of H3K9Me2 across the locus with corresponding loss of H3K27Ac in A38Lg, with no effect on A38Per (n=2 biological replicates, error bars=s.e.m.). RNAi against both CDH2 and TOP2B selectively blocked tumor formation in distant metastatic and precursor subclones that over-expressed these genes by RNA-seq, with no effect on A38Per or HPDE cells (n=4 technical replicates for each, error bars=s.d.m., *p<0.01; scale bars: 200 μm). Specific RNAi knockdown of gene expression is shown.

[0194] As illustrated in FIGS. 7A-7C, reprogrammed chromatin across distant metastatic subclones was detected. Western blots showed minimal or inconsistent changes for the indicated histone modifications between local-regional PDAC samples, including A38Per (Per). In contrast, a panel of distant metastatic subclones showed recurrent changes in specific modifications, compared to A38Per. Reprogramming was also observed between primary tumor subclones (Pr1, Pr2) and the lung met from the same patient.

[0195] As illustrated in FIG. 8A-8E, the specificity of reprogrammed histone modifications was established. Ki67 stains showed similar cell cycle rates for peritoneal and the matched lung met grown in serum, and serum-free media (SFM) arrested growth. Serial cell counts for the indicated times confirmed equal growth rates and growth arrest in SFM. GO analysis on RNA-seq data from cells cultured in serum vs. SFM further confirmed growth arrest in SFM (lung data, peritoneal gave identical results). Western blots showed persistence of reprogrammed chromatin modifications in serum/proliferative (-) and SFM/growth arrested (+) cells. Treatment of the peritoneal subclone with PDAC chemotherapies (Gem: Gemcitabine, G+FU: Gemcitabine+5-Fluorouracil) did not induce loss of methylation or gain of acetylation as seen between peritoneal and distant metastases, confirming that reprogramming was unrelated to treatment effects.

[0196] As illustrated in FIG. 9, an enrichment of heterochromatin modifications within LOCKs was identified. Plots of ChIP-seq read densities normalized to inputs for histone modifications (left labels) showed that heterochromatin modifications (H3K9Me2/3, H3K27Me3) were enriched in regions that were called LOCKs (0% to 100%, bottom panel

labels) for each subclone (indicated above graphs). In contrast, euchromatin modifications (H3K36Me3, H3K27Ac) were depleted from LOCKs.

[0197] As illustrated in FIGS. 10A-10B, the reprogramming of H3K9Me3 in LOCKs during PDAC subclonal evolution was observed. ChIP-seq data from paired peritoneal (A38Per) and lung (A38Lg) metastatic subclones detected dramatic reduction of H3K9Me3 in A38Lg, that overlapped with H3K9Me2 (which marks LOCK domains). Similar data for patient A13, which also showed loss of H3K9Me3 from LOCK regions in A13Pr2/A13Lg subclones, compared to the A1 3Pr1 primary tumor subclone.

[0198] As illustrated in FIGS. 11A-11B, the local reprogramming of DE gene loci within LOCKs was observed. Mapping DE genes from RNA-seq (distant mets and precursors vs. A38Per) to LOCKs revealed reciprocal changes in H3K27Me3 and H3K27Ac/H3K36Me3/DNA methylation around genes downregulated in LOCKs. The opposite changes in H3K27Me3 and H3K27Ac/H3K36Me3 were detected from genes upregulated in LOCKs. DNA methylation remained high in these regions. P-values for each comparison are listed in Supplementary Data 3.

[0199] As illustrated in FIG. 12, an enrichment of euchromatin modifications within ECDs was observed. Plots of ChIP-seq read densities normalized to inputs for histone modifications (left labels) showed that euchromatin modifications (H3K36Me3, H3K4Me3, H3K27Ac) were enriched in regions that were called ECDs (0% to 100%, bottom panel labels) for each subclone (indicated above the graphs). In contrast, heterochromatin modifications (H3K9Me2/3, H3K27Me3) were depleted from ECDs.

[0200] As illustrated in FIGS. 13A-13E, the reprogramming of large LOCKs during PDAC evolution was observed. H3K9Me3 was enriched and DNA hypomethylated in large LOCK domains. Striking reprogramming of H3K9Me3/2 and DNA methylation was detected in a subset of A38 large LOCKs. H3K9Me3 was also enriched across large HPDE LOCKs. Examples of reprogrammed domains between samples were obtained.

[0201] As illustrated in FIGS. 14A-14F, malignant heterogeneity between A38 subclones was discovered. Oxidoreductase capacity was measured with MTT assays performed on equal numbers of growth-arrested cells in the absence of serum, and MTT signals normalized to total cell numbers per well. Consistent with GO results, A38Lg possessed higher oxidoreductase activity. n=4 technical replicates. NADPH/NADP levels were measured with enzyme cycling assays on equal numbers of growth arrested cells. More NADPH/million cells were detected in A38Lg. n=2 biological replicates. A38Per and A38Lg maintained well/poorly differentiated morphology in patient tissues and across three separate in vitro culture conditions as indicated. IF performed on fixed tissues from the primary tumor showed loss of E-cadherin with gain of vimentin in the precursor subclone that seeded A38Lg, consistent with EMT. RNAi knockdown of KRAS blocked 3-D tumor formation in suspension assays more efficiently in A38Per than A38Lg. n=4 technical reps.

[0202] As illustrated in FIG. 15A-15C, it was found that rearrangements were targeted to Large LOCKs and ECDs. Total breakpoints were not significantly enriched within Large LOCKs or ECDs. Unlike typical LOCKs, Large LOCK/ECD breakpoints were significantly joined to breakpoints from homologous domains to form rearrangements.

Examples of Large LOCK rearrangements that generated translocations and amplifications were observed.

[0203] As illustrated in FIGS. 16A-16B, an enhanced glucose metabolism with depleted 6PG levels across distant metastases was observed. Extra-cellular glucose consumption and lactate secretion were elevated in distant mets relative to per. (n=3). Schematic of glycolytic (outside) and PPP (boxed) metabolites with intra-cellular metabolite levels plotted for each sample. Data represent LC-MS signals normalized to protein (n=3-5).

[0204] As illustrated in FIGS. 17A-17C, it was found that 6AN targets glucose metabolism and the PGD step of the PPP. 6AN selectively slowed rates of extra-cellular glucose consumption and lactate secretion in metastatic subclones with no effect on glutamine/glutamate. 6AN reduced incorporation of intracellular ⁰¹³-labeled glucose into metabolites downstream of the PPP. 6AN greatly increased metabolite levels of PGD substrate (6PG) and upstream metabolites (G1,5L) with corresponding reductions in downstream products.

[0205] As illustrated in FIGS. 18A-18C, it was found that 6AN selectively modulated the reprogrammed chromatin state of distant metastatic subclones. 6AN treatments generally increased global H3K9Me2 with corresponding decreased acetylation in distant metastatic subclones. Normal cells and local-regional PDACs did not show such changes. 6AN changes persisted after 3d treatment (+) followed by removal of 6AN from the media (+-) for an additional 3d.

[0206] As illustrated in FIGS. 19A-19D, it was found that 6AN targeted reprogrammed LOCK regions. Mapping 6AN repressed DE genes to A38Lg LOCKs revealed that these were located in reprogrammed LOCK-EI regions. ChIP-seq on DMSO vs. 6AN treated A38Lg detected a quantitative increase in LOCK-wide H3K9Me2 from reprogrammed regions (as aligned to A38Per LOCKs). ChIP-seq also detected 6AN-reduced H3K27Ac specifically from genes repressed in LOCKs with unchanged H3K27Me3.

[0207] As illustrated in FIGS. 20A-20C: 6AN selectively blocked tumor formation in distant metastatic subclones. 6AN did not interfere with the ability of local-regional PDAC samples to form tumors in 3-D Matrigel® assays or in 3-D suspension tumorsphere assays. n=2-4. In contrast, 6AN strongly blocked the ability of distant metastatic subclones to form tumors in 3-D suspension tumorsphere assays (shown) and 3-D Matrigel® assays (FIG. 5B). n=4, p<0.003. Scalebars: 200 uM.

[0208] As illustrated in FIGS. 21A-21C, the reprogramming of the TOP2B locus in response to 6AN was observed. RT-qPCR (top panels) showed that 6AN selectively repressed TOP2B the lung metastatic subclone. Similarly, ChIP assays showed that 6AN induced spreading of H3K9Me2 across the locus in the lung metastasis, with no effect on the paired peritoneal subclone. Representative RT-qPCR verified RNAi knockdown of CDH2 and TOP2B with minimal effect on vimentin (normalized to ERK, which was equally expressed across all conditions).

Discussion

[0209] The first major result of this study was widespread epigenetic reprogramming during the evolution of distant metastasis in the absence of metastasis-specific driver mutations, i.e., those not already present in the founder clone of the primary tumor. These involved large-scale reprogram-

ming of histone H3K9 and DNA methylation within large heterochromatin domains (LOCKS and hypomethylated blocks), as well as regional changes in gene regulatory modifications (H3K27Ac, H3K36Me3). Second, these changes specified heterogeneous malignant properties that emerged during subclonal evolution. In particular, evolutionarily divergent subclones from the same patient showed changes in gene expression from reprogrammed regions consistent with their individual malignant properties, including oxidoreductase capacity, differentiation state, chemoresistance, oncogene addiction, and patterns of genome instability. Third, it was the PGD step of the oxPPP that controlled aspects of reprogrammed chromatin and tumorigenicity in distant metastatic subclones, as shown by metabolomics, genetic knockdown of PGD, chemical inhibition of PGD, and knockdown of downstream target genes. This strongly suggests that this anabolic glucose pathway was selected during the evolution of distant metastasis to maintain malignant epigenetic state and tumorigenic properties.

[0210] These findings also raise several important but complex questions, which we are pursuing in other studies. Perhaps the most complicated pertains to the extent of epigenetic and malignant heterogeneity between subclones across patients. Just to answer this in a single patient, a combination of whole-genome mapping, RNA-seq, bioinformatics, and several downstream experimental approaches were required. We hypothesize that such heterogeneity is a function of evolutionary time: patients who present with late-stage, widely metastatic disease may possess more epigenetic and malignant divergence between subclones in their tumors than patients who present with early-stage disease. This possibility underscores the pressing need to detect cancers early, before such malignant heterogeneity arises.

[0211] Also unclear are the precise mechanisms whereby PGD/oxPPP activity controls global epigenetic state, which are likely to be complex. This could be mediated through any of the known oxPPP-dependent changes in cellular metabolism, including redox balance, fatty acid biosynthesis, and/or ribose biosynthesis, any of which can affect global epigenetic state through control of metabolite cofactors that activate or inhibit entire classes of chromatin modifying enzymes. PGD activity itself is also complex and subject to several modes of regulation, including transcriptional over-expression, post-transcriptional repression, post-translational modification, protein:protein interactions, substrate availability, feedback inhibition, cross-talk with other pathways, and subcellular localization including a highly conserved yet uncharacterized nuclear fraction (C. Lyssiotis, personal communication). PGD-dependence may be selected for by any of these mechanisms during the evolution of distant metastasis in different patients.

[0212] A final question is how global epigenetic changes are targeted to specific chromatin domains that encode gene expression changes during subclonal evolution. We hypothesize that transcription factors and chromatin modifying enzymes that directly bind these regions play major roles in targeting the reprogramming events, and several candidates were recurrently over-expressed in our RNA-seq datasets. This includes the histone demethylase KDM1A (LSD1), which could be particularly important since we previously

showed that this enzyme controls LOCK reprogramming and other studies have shown that it regulates breast cancer metastasis.

[0213] In summary, our findings in conjunction with deep sequencing studies on many of the same samples reported here suggest a model whereby driver mutations arise early to initiate PDAC tumorigenesis, followed by a period of subclonal evolution that generates heterogeneous metabolic, epigenetic, and malignant properties. Like driver mutations, those properties that confer increased fitness to cells that acquire them may be selected for and clonally expanded during invasive tumor growth and metastatic spread. The strong oxPPP-PGD dependence we observe in distant metastatic subclones could reflect such selection: distant metastatic sites provide ample glucose to fuel the pathway, pathway products (glucose-dependent NADPH) reduce oxygen species encountered within the sites, and the pathway itself is coupled to epigenetic programs that promote tumorigenesis. As such, reversal of malignant epigenetic programs by targeting the oxPPP could represent an effective therapeutic strategy for metastatic PDAC, one of the most lethal of all human malignancies.

Data Deposits

[0214] All ChIP-seq, RNA-seq, and bisulfite-seq sequencing data has been deposited online (GEO Number: GSE63126) at the following URL: ncbi.nlm.nih.gov/geo/query/acc.cgi?token=sxyjkaqsvfalheh&acc=GSE63126

Methods Summary

[0215] Tissue samples and cell lines were previously collected from PDAC patients by rapid autopsy, sequenced-validated, and monitored for mycoplasma as previously described. Low passage (2-17) rapid autopsy cell lines were cultured at 37° C. in DMEM with 10% fetal bovine serum (FBS, Gibco). For MTT assays, 15,000 cells/well were plated into 96 well plates in triplicate, treated 12 hours later, and assayed after 24 hr (glucose responses) or 6 days (chemotherapy) with CellTiter96 (Promega). For glucose response assays, nutrient-deplete DMEM (no glucose, glutamine, pyruvate, or serum) was used with addition of glucose as indicated. For glucose-dependent oxidative stress analysis, cells plated in triplicate and grown to 80% confluence followed by incubation in nutrient-deplete DMEM containing 10% dialyzed FBS with or without 10 mM glucose and 1 mM H₂O₂ for 24 hours. Cells were then washed with PBS, trypsinized, and viable cells counted with a hemocytometer. Glucose uptake and lactate secretion were measured with a YSI 7100 Bioanalyzer as described in the supplementary methods. For ¹³C-1,2 glucose tracing and steady state metabolite profiling, the Q Exactive MS (QE-MS; Thermo Scientific) coupled to liquid chromatography (LC Ultimate 3000 UHPLC) was used for metabolite separation and detection as previously described. Detailed conditions are provided in the supplementary methods.

[0216] Histones were acid extracted as described and western blots performed on 3.5 ug histones, which were checked by Ponceau stains prior to western blot to ensure equal loading. Densitometry was performed with ImageJ software. RNA was extracted with Trizol reagent (Life Technologies) and isopropanol precipitated. Genomic DNA was purified with MasterPure DNA extraction reagents (Epicenter). Immunohistochemistry, H&E staining, and

immunofluorescence on formalin-fixed, paraffin-embedded (FFPE) tissue microarray sections (TMAs) were performed according to standard procedures. Antibodies used for western blot, IHC, and ChIP are listed in Table 11.

factor receptor mediates increased cell proliferation, migration, and aggregation in esophageal keratinocytes in vitro and in vivo. *J Biol Chem* 278, 1824-30 (2003)). Briefly, 6 well permeable transwell plates (Costar 3414) were over-

TABLE 11

Antibodies and conditions used for western blots, immunostain, and ChIP experiments				
Antibody	Source	Catalogue #	Western dilution	Immuno dilution
CDH1	Cell Signaling	4065	1/500	1/100, IF
Vimentin	NeoMarkers	Ms-129-P	1/500	1/100, IF
G6PD	Cell Signaling	8866S	1/500	N/A
PGD	Cell Signaling	13389S	1/500	N/A
TKT	Cell Signaling	8616S	1/500	N/A
TALDO1	Santa Cruz	Sc-134795	2 ug/ml	N/A
ERK1/2	Cell Signaling	4696S	1/500	N/A
p-ERK1/2	Cell Signaling	4370S	1/500	N/A
KRAS	Santa Cruz	Sc-30	2 ug/ml	N/A
RPE	Santa Cruz	Sc-162124	2 ug/ml	N/A
RPIA	Abcam	Ab181235	1/500	N/A
CD44	Cell Signaling	5640S	1/500	N/A
Epcam	Millipore	CBL251	1/500	N/A
CDH2	Cell Signaling	4061S	1/250	N/A
H3K9Me2	Abcam	ab1220	1/10,000	1/5000 IHC, IF
H3K9Me3	Abcam	ab8898	1/10,000	1/5000 IHC
H3K27Me3	Millipore	07-449	1/7500	N/A
H3K9Ac	Millipore	07-352	1/10,000	N/A
H3K36Me3	Abcam	ab9050	1/10,000	1/5000 IHC
H3K27Ac	Abcam	ab4729	1/5000	N/A
H4K20Me3	Millipore	07-463	1/1000	1/5000 IHC
H4K16Ac	Millipore, Abcam	07-329, ab109463	1/10,000	N/A
total H3	Abcam	ab 1791	1/30,000	N/A
total H4	Abcam	ab 10158	1/30,000	N/A
YH2AX	Abcam	ab11174	1/5,000	N/A

[0217] RNAi experiments were performed with siRNA transfections (Oligofectamine, Life Technologies) using negative control siRNA (Sigma, SIC001) and pre-designed siRNA oligonucleotides against indicated genes in parallel (Sigma, PGD: SASI_Hs02_00334150, CDH2: SASI_Hs01_00153995, TOP2B: SASI_Hs02_00311874). siRNAs against mutant KRAS^{G12V} (CUACGCCAACAGCUC-CAAC) (SEQ ID NO:1) were custom designed. Cells were incubated with siRNAs for 4 days after transfection and harvested. For drug treatments in 2-D, cells were grown to 70-80% confluency and treated for 3 days with 250 uM 6AN or DMSO negative control.

[0218] 3-D matrigel assays were adapted from Cheung et al. (Control of alveolar differentiation by the lineage transcription factors GATA6 and HOPX inhibits lung adenocarcinoma metastasis. *Cancer Cell* 23, 725-38 (2013)). Briefly, 2-D cultures were trypsinized into single cells, 4,000 cells/mL were suspended and thoroughly mixed in ice-cold DMEM containing 5% matrigel (BD systems) and 2% FBS (+/-DMSO/6AN as needed), 500 ul plated in quadruplicate into 24 well ultra-low attachment plates and incubated for at least 7 days to allow tumor growth. Well-formed tumors were then counted and representative photographs taken with an EVOS instrument. 3-D suspension tumorsphere assays were performed with 20,000 starting cells/well in ultra-low attachment 6 well plates as described, and tumors counted/photographed after at least 7 days of tumor growth.

[0219] Organotypic tumor forming assays were adapted from Ridky et al. (P. A. Invasive three-dimensional organotypic neoplasia from multiple normal human epithelia. *Nat Med* 16, 1450-5 (2010)) and Andl et al. (Epidermal growth

layed with 1 mL type 1 collagen containing 10xDMEM (acellular layer). Human dermal fibroblasts (ATCC) were suspended (12×10^6 cells/mL) in a mixture of ice cold 10xDMEM, 10% FBS, 52.5% collagen, and 17.5% matrigel (cellular layer), thoroughly mixed, and 2 mL/well plated over the acellular layer. The mixture was allowed to partially solidify for approximately 15 minutes at 37° C., followed by triplicate injection of 1×10^6 PDAC (suspended in 20 ul DMEM) cells into the cellular layer. Cells were incubated for 24 hours in fibroblast growth media above and below the inserts to initiate contraction of the discs. Fresh media with DMSO or 6AN was then added and replenished every 2 days for 6 additional days, followed by addition of DMEM with DMSO or 6AN underneath the inserts (no media on the top) for an additional 7 days. Discs were harvested, fixed overnight in 10% formalin, thinly sectioned, paraffin embedded, and stained with H&E. Tumors were photographed and measured with an Olympus BX53 microscope using cellSens Standard software.

[0220] Tests for statistical significance (two-tailed students t-test) were performed on data collected from technical replicate (performed in parallel at the same time) or biological replicate (performed at different times) experiments as indicated in the figure legends using excel software for western blot densitometry, MTT assays, and tumor measurements. Whole genome bisulfate sequencing and RNA-seq were performed with HiSeq instruments (Illumina) as described in Hansen et al. (Increased methylation variation in epigenetic domains across cancer types. *Nat Genet* 43, 768-75 (2011)). ChIP assays were performed as previously described for fixed cells with sonication. For ChIP-qPCR,

equal amounts of paired input/IP DNA were amplified by real-time PCR (Roche LightCycler96) and fold enrichments calculated. Primer sequences are listed in Supplementary Table 11. For ChIP-seq, immunoprecipitated and input DNA was further sheared to 200-300 bp fragments, size-selected on agarose gels, and sequenced on either HiSeq (Illumina) or SOLiD (Applied Biosystems) formats with comparable results. IP sequencing reads were normalized to their corresponding inputs. Sequencing procedures, bioinformatics methods including domain calls, and statistical analyses are described in detail within the supplementary methods section.

TABLE 12

Real-time PCR primer sequences used for ChIP-qPCR and RT-PCR experiments				
Locus/Assay Primer Name: Sequence (5'-3')			SEQ ID NO:	
CDH2 ChIP	Chr18_24.258F:	GCTCAGCCCTGTATCAGCCAGC	2	
CDH2 ChIP	Chr18_24.258R:	GGGTTACAGGTATGAGCCACTGC	3	
CDH2 ChIP	Chr18_24.506F:	AATGGAGAAGTCAGGAATGTAGTCC	4	
CDH2 ChIP	Chr18_24.506R:	GTATTTTCATTTATCAAGTTGCAGCTCC	5	
CDH2 ChIP	Chr18_24.834F:	TTTGCTTCTCACTCCAAGTTCATCC	6	
CDH2 ChIP	Chr18_24.834R:	CAACCTCAGGAACAATGCATCAGC	7	
CDH2 ChIP	Chr18_25.125.6F:	CGAAACAGTCCAGCTGCTATGG	8	
CDH2 ChIP	Chr18_25.125.6R:	CTTGGCTATTGTGACTGGTACTGC	9	
CDH2 ChIP	Chr18_25.428.000F:	CCAATGCACTAATTTAATGTCATGC	10	
CDH2 ChIP	Chr18_25.428.000R:	CGTGCTAATTTCTATGGTACACTGG	11	
CDH2 ChIP	Chr18_25.632F:	CCTAATCCAATATGCCTGGTGTCC	12	
CDH2 ChIP	Chr18_25.632R:	CTGGAAGTCTGAGATCAAGGTGC	13	
CDH2 ChIP	Chr18_25.778F:	AATAATCACGAAGCACTTCTGTATTGC	14	
CDH2 ChIP	Chr18_25.778R:	TCACCAGCAGACATAGTCATACTTCC	15	
CDH2 ChIP	Chr18_25.808F:	CCTTGGAGGTGGAGTCTACAGAGG	16	
CDH2 ChIP	Chr18_25.808R:	CTGCTAGCGTAGCCATCTGAGATCG	17	
TOP2B ChIP	Chr3_25.398F:	GCCCTGTCTTCCCAGAATCATTGC	18	
TOP2B ChIP	Chr3_25.398R:	CATGAAGCCTATGAAGATCATTATGG	19	
TOP2B ChIP	Chr3_25.540F:	TTTAGCCAGCAAGTATTCTAGCATGG	20	
TOP2B ChIP	Chr3_25.540R:	GTCAGTGTGATTCAAGTAACAATGATGG	21	
TOP2B ChIP	Chr3_25.622F:	CCTGCTCAAGGCTGACATGTCACC	22	
TOP2B ChIP	Chr3_25.622R:	GTCGGACTCGATGGTCAGCACTGG	23	
TOP2B ChIP	Chr3_25.733F:	AACCCGAACTTTCAATGCACTTGG	24	
TOP2B ChIP	Chr3_25.733R:	CTTCCTCTATAGTGAAGACCCTAGG	25	
TOP2B ChIP	Chr3_25.812F:	TATGGCCATTCTTGCAGCAGTAAGG	26	
TOP2B ChIP	Chr3_25.812R:	AAAGTTGGCTAAGGACATGAATAGGC	27	
TOP2B ChIP	Chr3_25.973F:	GGAGATTCCCTCAGGTGCCTATACC	28	
TOP2B ChIP	Chr3_25.973R:	CTGGTGTTCCAGGCACCACTGAGG	29	
CDH2 RT-PCR	CDH2F:	TTATTACTCCTGGTGCAGT	30	
CDH2 RT-PCR	CDH2R:	GAGCTGATGACAAATAGCGG	31	
TOP2B RT-	TOP2BF:	GTTACAGGTGGTCGTAATGGTT	32	
TOP2B RT-	TOP2BR:	TTGGCTTCAGAAGTCTTCATCA	33	

Supplementary Methods

[0221] YSI metabolite analysis. Metabolite consumption (glucose and glutamine) and production (lactate and glutamate) were measured using a YSI 7100 Bioanalyzer. Indicated cell lines were plated at day -1 in a 6-well plate. At day 0 cells were counted (3 wells) or cultured in either regular medium or medium supplemented with the indicated compound. Tissue culture supernatants (1 mL, n=3, each condition) were harvested 72 hours after cell plating. Tissue culture conditions were optimized to ensure nutrient availability and exponential cell growth. Metabolite consumption/production data were normalized to cell number area under the curve, as previously described (Lee et al 2014: PMID: 24998913). The area under the curve (AUC) was calculated as $N(T)d/\ln 2(1-2^{-T/d})$, where N(T) is the final cell count, d is doubling time, and T is time of experiment. Doubling time was calculated as $d=(T)[\log(2)/\log(Q2/Q1)]$, where Q1 is starting cell number and Q2 is final cell number, as determined by manual counting using a hemocytometer.

[0222] LC-HRMS Metabolite Profiling. LC-HRMS samples were prepared and analyzed as described in Liu et al. (Development and quantitative evaluation of a high-resolution metabolomics technology. *Anal Chem* 86, 2175-84 (2014)). For glucose tracing experiments, cells were plated into 6 well plates in triplicate, grown in DMEM with 10% FBS until 70-80% confluent, washed 2x with nutrient deplete DMEM, and incubated in nutrient deplete DMEM containing 10 mM ^{13}C -1,2 labeled glucose (Cayman) and 10% dialyzed FBS (Invitrogen) for an additional 36 hours. Additional replicates were also included and counted at the end of the experiment for normalization. Metabolism was quenched by quickly removing media and adding 1 mL pre-chilled (-80°C .) LC-MS grade 80% methanol (Sigma), incubated at -80°C . for at least 20 minutes, followed by scraping into the methanol and pelleting of metabolites by centrifugation. For drug treatments, cells were incubated in standard DMEM+/-DMSO/6AN for 36 hours, followed by incubation in labeled glucose media+/-DMSO/6AN for an additional 36 hours, quenched and pelleted as above. Pellets were reconstituted in equal volumes of 1:1 LC-MS grade acetonitrile:methanol and water and 5 μL were injected to the LC-QE-MS for analysis. For steady state measurements cells were incubated in growth media (DMEM with 10% FBS for PDACs, keratinocyte serum-free media for HPDE) until they reached 80-90% confluence, followed by 48 hours in DMEM without serum (for PDACs, since the standard growth media for comparison HPDE cells also did not contain serum). Metabolism was then quenched with methanol and metabolites pelleted as above. Pellets were reconstituted into a volume normalized to protein content (15 μL of 1:1 acetonitrile:methanol and 15 μL of water was used per 1 mg protein) and analyzed by LC-QE-MS. Raw data collected from the LC-QE-MS was processed on Sieve 2.0 (Thermo Scientific) using a targeted frame-seed that included glycolytic/PPP metabolites as required for the analysis. The output file including detected m/z and relative intensity in different samples is obtained after data processing, and replicates of selected metabolites from each sample were graphed and presented as shown in the figures.

[0223] Preparation of sequencing libraries. Libraries were prepared from 2-10 ng of IP ChIP DNA and 100 ng of input DNA and sequenced on Illumina HiSeq (APF laboratory). Briefly, samples were checked for quality and concentration from 150-250 bp on a bioanalyzer. DNA was end-repaired

using Klenow polymerase in 58 μL of reaction buffer. For IP DNA, Klenow was diluted 1:5. Samples were incubated at 20°C . for 30 minutes and subsequently purified on QIAquick PCR purification columns. A-tails were then added to the DNA with Klenow and dATP in NEB buffer 2 at 37°C . for 30 minutes and cleaned with Qiagen MiniElute PCR purification columns. Sequencing adapters were then ligated onto the DNA for 15 minutes at room temperature followed by cleaning with MiniElute columns. Samples were then run on 2% agarose gels and DNA from 216 bp-366 bp (DNA plus adapters) were cut from the gel and purified with Qiagen Gel extraction kits. Concentrations were then checked on a bioanalyzer and 8 ng were PCR amplified with Phusion polymerase (Fisher) for 15 cycles (10 sec 98°C ., 30 sec 65°C ., 30 sec 72°C .) followed by 5 minutes at 72°C . Samples were then cleaned with Ampure kits (Illumina) and washed with 80% ethanol. DNA samples were resuspended at the end of the cleanup into 17.5 μL buffer EB (Qiagen) and subjected to next generation sequencing on Illumina HiSeq platform according to manufacturer's instructions. For SOLID sequencing, ChIP DNA was prepared and samples were processed according to manufacturer's protocols in the Johns Hopkins CRBII core facility.

[0224] BS-Seq data processing. 100 bp paired-end HiSeq2000 sequencing reads were aligned by BSmooth bisulfite alignment pipeline (version 0.7.1) as previously described in Hansen et al. (Increased methylation variation in epigenetic domains across cancer types. *Nat Genet* 43, 768-75 (2011)). Briefly, reads were aligned by Bowtie2 (version 2.0.1) against human genome (hg19) as well as the lambda phage genome. After alignment, methylation measurements for each CpG were extracted from aligned reads. We filtered out measurements with mapping quality <20 or nucleotide base quality on cytosine position <10 and we also removed measurements from the 5' most 10 nucleotides of both mates. Then, bsseq package in BSmooth was used to identify small and large differentially methylated regions (DMRs). Only CpGs with at least coverage of 3 in all samples were included in our analysis. For small DMRs, smooth window of 20 CpGs or 1 kb was used, and t-statistic cutoff of -4.6, 4.6 and methylation difference greater than 20% were used for identifying small DMRs. While for large DMRs, smooth window of 200 CpGs or 10,000 bps was used, and t-statistic cutoff of -2, 2, methylation difference greater than 10% and length of DMRs >5 kb were used for identifying large DMRs.

[0225] RNA-Seq data processing. 100 bp paired-end HiSeq2000 sequencing reads were aligned against human genome (hg19) by OSA (version 2.0.1) with default parameters. After alignment, only uniquely aligned reads were kept for further analysis. Gene annotation information was downloaded from ENSEMBL (ensembl.com, release 66). Reads count for each gene of all samples were estimated using HTSeq (huber.embl.de/users/anders/HTSeq/doc/overview.html) and then were used to identify differentially expressed (DE) genes using DESeq package. Genes with FDR <0.01 and fold-change >1.5 were considered DE genes.

[0226] Chip-seq data processing. For 46 bp paired-end Illumina HiSeq2000 sequencing data, reads were aligned against human genome (hg19) using BWA with default parameters as described in Li et al. (Fast and accurate short read alignment with Burrows-Wheeler transform. *Bioinformatics* 25, 1754-60 (2009)). After alignment, duplicate reads

were removed and only uniquely aligned reads were kept for further analysis. For 48 bp single-end Solid sequencing data, reads were aligned using Bowtie⁷ with default parameters and only uniquely aligned reads were kept for further analysis. For narrow histone modification peaks (H3K4Me3 and H3K27Ac), MACS2 were used for peak calling with default parameters⁸. For broad histone modification enrichments (H3K36Me3, H3K27me3, and H3K9Me2/3), peak calling was performed using RSEG which is based on hidden Markov model (HMM) and specifically designed for identifying broad histone peaks (see Song et al. (Identifying dispersed epigenomic domains from ChIP-Seq data. *Bioinformatics* 27, 870-1 (2011))).

[0227] Identifying large chromatin domains. We define LOCK domains for heterochromatin modifications (H3K9Me2/H3K27Me3) based on the peak calling results from RSEG. Briefly, peaks shorter than 5 kb were first removed to prevent regions with many nearby, short peaks being called as LOCKs. Then, neighboring peaks with distance less than 20 kb were merged to into one domain. Merged regions greater than 100 Kb identified in both biological replicates were called LOCKs. We noticed another unique subset of LOCKs that were invariably larger than 500 kb, strongly enriched with H3K9Me3, depleted of H3K9Me2 and H3K27Me3, and flanked by strong peaks of H3K27Me3 at the boundaries. Because of this, we defined these LOCKS by H3K9Me3 regions with length greater than 500 kb and less than 50% of their length overlapped with H3K27me3. Finally, the large regions (>50 kb) between heterochromatin domains that contained at least one gene with corresponding euchromatic H3K4Me3/H3K27Ac regulatory peaks were defined as ECDs. Because we found that H3K27Ac alone was sufficient for these calls, H3K4Me3 was also used for the initial test dataset with A38Per/Lg, but not required in subsequent datasets (A13Pr1/2, A13Lg). All codes used for domain calls are available upon request.

[0228] Defining different gene groups. Genes were classified as belonging within euchromatin (>50% of genic region located in ECDs) or heterochromatin (>50% of genic region located in those heterochromatin domains including LOCKS and G-LOCKS). A handful of other genes that did not fit these criteria and were classified as “other”.

[0229] Quantifying and enrichment plotting of ChIP-seq and RNA-seq. To plot each histone modification on defined large chromatin domains and their flanking regions, we divided flanking sequences of chromatin domains into bins with fixed length (in bp) and domains themselves into bins with fixed percentage of each domain length. ChIP enrichment was measured and normalized as described in Hawkins et al. (Distinct epigenomic landscapes of pluripotent and lineage-committed human cells. *Cell Stem Cell* 6, 479-91 (2010)). In brief, the number of reads per kilobase of bin per million reads sequenced was calculated for each ChIP and its input control (denoted as $RPKM_{chip}$ and $RPKM_{input}$). ChIP enrichment is measured as $\Delta RPKM = RPKM_{ChIP} - RPKM_{input}$ and ChIP enrichment regions should have $\Delta RPKM > 0$. Then all $\Delta RPKM$ were normalized to a scale between 0 and 1 and the average normalized ChIP enrichment signals across all large chromatin domains were plotted for each histone mark. RNA-Seq data was also normalized by the number of reads per kilobase of bin per million reads sequenced and plotted similarly.

Supplementary Data

[0230] Supplementary Data 1. This lists numbers of sequencing reads (total reads and uniquely aligned reads) for all replicate samples for ChIP-seq, WGBS, and RNA-seq experiments and includes correlation coefficients between the replicate samples.

[0231] Supplementary Data 1A: Summary of ChIP-seq reads for all replicate samples

Samples (Name_replicate#_IP antibody)	Total reads	Uniquely aligned reads
38Per_1_K27ac	27,100,820	23,396,028
38Per_2_K27ac	20,750,076	17,913,278
38Lg_1_K27ac	22,821,996	19,946,514
38Lg_2_K27ac	25,996,970	22,475,058
38Per_1_K9ac	27,148,594	22,200,344
38Per_2_K9ac	25,585,898	18,706,456
38Lg_1_K9ac	26,607,612	21,750,232
38Lg_2_K9ac	28,316,524	23,222,230
38Per_1_K4me3	28,059,734	23,718,542
38Per_1_K4me3	43,068,697	20,704,383
38Lg_2_K4me3	24,455,456	21,184,788
38Lg_2_K4me3	44,715,633	17,458,603
38Per_1_K36me3	44,738,783	23,961,134
38Per_2_K36me3	25,140,434	19,239,814
38Lg_1_K36me3	25,532,398	18,263,078
38Lg_2_K36me3	48,929,336	24,747,992
38Per_1_K27me3	26,546,612	21,523,318
38Per_1_K27me3	47,076,156	21,613,912
38Lg_2_K27me3	25,528,442	20,886,386
38Lg_2_K27me3	44,444,667	16,568,247
38Per_1_K9me2	69,629,048	55,830,176
38Per_K9me2_2	64,949,506	51,281,848
38Lg_K9me2_1	51,787,694	39,725,736
38Lg_2_K9me2	71,421,136	55,643,850
38Per_1_K9me3	28,530,078	18,843,038
38Per_2_K9me3	30,869,994	19,860,544
38Lg_1_K9me3	33,055,178	20,251,534

-continued

Samples (Name__replicate#__IP antibody)	Total reads	Uniquely aligned reads
38Lg__2__K9me3	36,205,418	22,568,940
38Per__1__Input	28,748,102	22,637,100 for 38Per__1 K16ac, K27ac, K9ac, K4me3, K27me3, K9me2 and K9me3
38Per__1__Input	43,698,893	20,955,594 for 38Per__1 Solid K4me3, K36me3 and K27me3
38Per__1__Input	78,376,368	61,062,024 for 38Per__1 K9me2
38Per__2__Input	26,162,800	20,358,392 for 38Per__2 K16ac, K9ac, K36me3, K9me2 and K9me3
38Per__2__Input	22,795,752	17,717,432 for 38Per__2 K27ac
38Per__2__Input	73,425,982	57,237,686 for 38Per__2 K9me2
38Lg__1__Input	30,391,444	23,128,054 for 38Lg__1 K16ac, K27ac, K9ac and K9me3
38Lg__1__Input	26,527,468	20,878,592 for 38Lg__1 K36me3 and K9me2
38Lg__1__Input	56,918,912	44,006,646 for 38Lg__1 K9me2
38Lg__2__Input	24,372,670	21,204,480 for 38Lg__2 K16ac, K27ac, K9ac, K4me3, K27me3 and K9me3
38Lg__2__Input	43,698,893	19,884,219 for 38Lg__2 Solid K4me3 and K27me3
38Lg__2__Input	42,655,553	25,391,128 for 38Lg__2 Solid K36me3
38Lg__2__Input	82,576,576	65,151,312 for 38Lg__2 K9me2
13Pr2__1__K27Me3	45,562,122	35,405,382
13Pr2__2__K27Me3	48,647,198	37,601,820
13Pr2__1__K36Me3	50,525,636	40,320,884
13Pr2__2__K36Me3	43,172,826	34,777,556
13Pr2__1__K9Me3	52,034,354	28,398,244
13Pr2__2__K9Me3	47,846,284	27,025,984
13Pr2__1__K27Ac	39,057,664	32,754,504
13Pr2__2__K27Ac	45,988,666	38,419,042
13Pr2__1__K9Me2	36,367,760	27,747,040
13Pr2__2__K9Me2	49,421,058	36,685,524
13Pr2__1__Input	42,932,496	34,453,136 for 13Pr2__1 K27Ac and K9Me2
13Pr2__2__input	39,166,036	31,527,030 for 13Pr2__2 K27Ac and K9me2
13Pr2__1__Input	43,812,050	33,019,642 for 13Pr2__1 K36Me3
13Pr2__2__Input	42,483,790	32,000,834 for 13Pr2__2 K36Me3
13Pr2__1__Input	36,646,218	27,841,600 for 13Pr2__1 K27Me3 and K9Me3
13Pr2__2__Input	37,605,184	28,502,806 for 13Pr2__2 K27Me3 and K9Me3
13Pr1__1__K27Me3	42,464,788	32,291,936
13Pr1__2__K27Me3	42,717,990	32,886,274
13Pr1__1__K36Me3	45,628,144	34,403,256
13Pr1__2__K36Me3	47,010,260	35,169,858
13Pr1__1__K27Ac	42,969,352	36,362,084
13Pr1__2__K27Ac	52,912,056	43,709,782
13Pr1__1__K9Me3	43,261,454	27,946,136
13Pr1__2__K9Me3	48,134,638	30,111,834
13Pr1__1__K9Me2	51,237,440	37,126,410
13Pr1__2__K9Me2	44,260,552	34,079,972
13Pr1__1__K4Me3	47,817,718	40,460,332
13Pr1__2__K4Me3	39,916,242	34,219,430
13Pr1__1__Input	27,015,968	21,991,728 for 13Pr1__1 K9Me3 and K9Me2
13Pr1__2__Input	42,754,918	34,651,668 for 13Pr1__2 K9Me3 and K9Me2
13Pr1__1__Input	37,703,540	30,589,064 for 13Pr1__1 K36Me3 and K27Ac
13Pr1__2__Input	37,251,906	30,188,840 for 13Pr1__2 K36Me3 and K27Ac
13Pr1__1__Input	45,330,014	36,495,018 for 13Pr1__1 K27Me3 and K4Me3
13Pr1__2__Input	41,671,660	33,507,094 for 13Pr1__2 K27Me3 and K4Me3
13Lg__1__K27Me3	46,174,838	35,837,518
13Lg__2__K27Me3	47,763,132	36,963,900
13Lg__1__K36Me3	48,735,296	39,047,890
13Lg__2__K36Me3	44,439,570	35,973,060
13Lg__1__K27Ac	52,682,716	44,187,348
13Lg__2__K27Ac	38,043,964	32,298,344
13Lg__1__K9Me3	44,449,474	25,717,050
13Lg__2__K9Me3	49,020,848	28,456,692
13Lg__1__K9Me2	40,580,872	31,979,380
13Lg__2__K9Me2	42,760,754	33,395,094
13Lg__1__input	53,086,242	42,631,966 for 13Lg__1 K27Ac and K9Me2
13Lg__2__input	41,676,088	33,503,572 for 13Lg__2 K27Ac and K9Me2
13Lg__1__Input	40,822,392	31,354,916 for 13Lg__1 K36Me3
13Lg__2__Input	45,292,342	35,189,166 for 13Lg__2 K36Me3
13Lg__1__Input	49,243,126	37,741,490 for 13Lg__1 K27Me3 and K9Me3
13Lg__2__Input	47,226,322	36,317,110 for 13Lg__2 K27Me3 and K9Me3
HPDE__1__K27Me3	41,619,730	1,644,221,182
HPDE__2__K27Me3	45,844,562	1,870,133,582
HPDE__1__K36Me3	38,327,244	28,084,826

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Samples (Name__replicate#__IP antibody)	Total reads	Uniquely aligned reads
HPDE__2__K36Me3	35,519,752	27,012,400
HPDE__1__K27Ac	52,294,724	43,428,888
HPDE__2__K27Ac	35,271,886	30,001,598
HPDE__1__K9Me3	49,006,354	31,150,780
HPDE__2__K9Me3	49,995,186	31,313,578
HPDE__1__K9Me2	47,415,884	36,297,824
HPDE__2__K9Me2	47,621,364	36,772,334
HPDE__1__K4Me3	45,262,500	39,528,934
HPDE__2__K4Me3	34,511,978	30,181,046
HPDE__1__Input	50,286,666	40,743,792 for HPDE__1__K9Me3 and K9Me2
HPDE__2__Input	45,780,736	37,154,676 for HPDE__2__K9Me3 and K9Me2
HPDE__1__Input	36,424,754	29,600,008 for HPDE__1__K36Me3 and K27Ac
HPDE__2__Input	48,330,362	39,054,926 for HPDE__2__K36Me3 and K27Ac
HPDE__1__Input	38,269,792	30,829,930 for HPDE__1__K27Me3 and K4Me3
HPDE__2__Input	37,635,368	30,421,766 for HPDE__2__K27Me3 and K4Me3
38Lg__DMSO__1__K27Me3	45,364,340	33,889,440
38Lg__DMSO__2__K27Me3	37,628,254	29,077,804
38-5__DMSO__1__K9Me2	57,024,354	42,891,924
38Lg__DMSO__1__K27Ac	44,665,878	37,664,038
38-5__DMSO__1__Input_batch4	69,705,942	56,571,128 for 38-5__DMSO__1__K9Me2
38Lg__DMSO__1__Input	37,296,752	30,470,440 for 38Lg__DMSO__1__K27Ac
38Lg__DMSO__1__Input	49,727,864	40,284,668 for 38Lg__DMSO__1__K27Me3
38Lg__DMSO__2__Input	40,505,458	32,886,902 for 38Lg__DMSO__2__K27Me3
38Lg__6AN__1__K27Me3	50,310,568	39,528,064
38Lg__6AN__2__K27Me3	38,884,546	32,325,978
38-5__6AN__1__K9Me2	33,324,396	24,956,330
38Lg__6AN__1__K27Ac	40,895,998	34,565,998
38-5__6AN__1__Input_batch4	42,480,878	34,537,520 for 38-5__6AN__1__K9Me2
38Lg__6AN__1__Input_batch3	40,615,920	33,033,908 for 38Lg__6AN__1__K27Ac
38Lg__6AN__1__Input_batch5	45,637,332	37,039,730 for 38Lg__6AN__1__K27Me3
38Lg__6AN__2__Input_batch5	36,771,474	29,835,218 for 38Lg__6AN__2__K27Me3
Total	7,441,856,062	

[0232] Supplementary Data 1B: Summary of WGBS reads for all replicate samples

Samples	Total reads	Aligned reads	Total CpGs	Aligned CpGs	Coverage per replicate
38Per rep1	281,674,084	234,515,645	28,217,448	24,736,894	5.74
38Per rep2	279,677,926	233,819,498	28,217,448	24,668,800	5.65
38Lg rep1	246,354,388	242,041,867	28,217,448	24,079,382	4.96
38Lg rep2	293,066,340	203,060,037	28,217,448	24,568,234	5.87
38-Lv rep1	481,671,728	394,675,778	28,217,448	25,699,745	12.06
38-Lv rep2	550,721,820	443,244,002	28,217,448	25,797,654	13.57
13Pr2 rep1	470,251,572	378,466,223	28,217,448	25,763,514	11.83
13Pr2 rep2	515,334,056	421,725,378	28,217,448	25,287,251	12.01
13Pr1 rep1	421,949,244	340,966,300	28,217,448	25,733,843	10.58
13Pr1 rep2	383,229,986	308,746,570	28,217,448	25,613,462	9.58
13Lg rep1	489,959,480	400,936,046	28,217,448	25,342,333	11.59
13Lg rep2	505,568,682	415,435,288	28,217,448	25,358,384	11.74
HPDE rep1	319,334,144	259,838,538	28,217,448	25,498,338	8.14
HPDE rep2	409,472,974	329,823,993	28,217,448	25,755,008	10.07
38Lg DMSO rep1	381,562,458	313,237,618	28,217,448	25,509,209	9.67
38Lg DMSO rep2	462,809,566	378,726,614	28,217,448	25,716,790	11.67
38Lg 6AN rep1	668,822,798	546,162,307	28,217,448	25,987,966	16.62
38Lg 6AN rep2	408,683,276	329,596,955	28,217,448	25,586,266	10.20
A124PerMet	388,402,336	313,722,746	28,217,448	25,551,642	7.39
A124Pr	348,071,524	281,407,466	28,217,448	25,424,592	6.73
Normal	335,262,856	264,003,665	28,217,448	25,423,437	6.37

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Samples	Total reads	Aligned reads	Total CpGs	Aligned CpGs	Coverage per replicate
A125LvMet2	299,092,468	241,196,683	28,217,448	24,124,822	5.55
A125LvMet1	256,878,816	210,595,185	28,217,448	24,190,459	5.09
A125Pr2	274,964,060	226,288,821	28,217,448	24,326,375	5.39
A125Pr1	338,097,576	277,039,476	28,217,448	25,344,423	6.55
Total		7,989,272,699			

[0233] Supplementary Data 1C: Summary of RNA-seq reads for all replicate samples

Samples	Total reads	Uniquely aligned reads	Genes with at least one read
38Per rep1	136,981,522	126,989,515	24,629
38Per rep2	209,196,426	194,285,575	25,767
38Per SFM rep1	131,806,836	120,661,977	23,750
38Per SFM rep2	170,418,406	157,947,694	25,204
38Lg rep1	157,009,238	146,313,205	22,347
38Lg rep2	126,740,896	118,431,884	21,849
38Lg SFM rep1	157,587,660	144,483,739	23,085
38Lg SFM rep2	136,421,728	124,124,770	22,387
38-Lv rep1	121,404,350	99,124,421	23,033
38-Lv rep2	201,919,638	160,412,909	24,203
13Pr2 rep1	189,720,644	163,161,693	25,011
13Pr2 rep2	154,397,366	132,746,732	24,122
13Pr1 rep1	194,020,194	152,395,660	25,013
13Pr1 rep2	159,731,916	124,045,393	24,300
13Lg rep1	258,937,646	222,764,245	25,530
13Lg rep2	104,375,982	89,301,099	24,908
HPDE rep1	142,777,652	115,147,712	23,274
HPDE rep2	148,752,028	119,731,612	23,348
38Per DMSO rep1	103,160,056	90,683,198	23,718
38Per DMSO rep2	205,086,938	175,208,405	25,285
38Per 6AN rep1	163,145,386	144,907,391	24,514
38Per 6AN rep2	163,346,616	145,684,561	24,049
38Lg DMSO rep1	183,580,116	158,065,524	23,609
38Lg DMSO rep2	177,808,970	156,924,338	23,334
38Lg 6AN rep1	231,471,388	201,726,950	24,544
38Lg 6AN rep2	181,401,400	157,469,565	23,376
Total		3,742,739,767	

[0234] Supplementary Data 1D: Summary of sequencing correlation coefficients for each replicate

Samples	Modification	Correlation coefficients between replicates
38Per	K4Me3	0.8487501
38Per	K36Me3	0.7908561
38Per	K27Me3	0.8087827
38Per	K9Me2	0.9533903
38Per	K9Me3	0.973981
38Per	K9Ac	0.8405352
38Per	K27Ac	0.9662306
38Per	K16Ac	0.8958976
38Lg	K4Me3	0.8977917
38Lg	K36Me3	0.7457856
38Lg	K27Me3	0.8283084
38Lg	K9Me2	0.9345982
38Lg	K9Me3	0.9870598
38Lg	K9Ac	0.8557192
38Lg	K27Ac	0.9749662

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Samples	Modification	Correlation coefficients between replicates
38Lg	K16Ac	0.8634484
13Pr2	K27Ac	0.9795735
13Pr2	K27Me3	0.957889
13Pr2	K36Me3	0.970403
13Pr2	K9Me2	0.9710885
13Pr2	K9Me3	0.9918936
13Pr1	K27Ac	0.9797524
13Pr1	K27Me3	0.9481441
13Pr1	K36Me3	0.9697859
13Pr1	K9Me2	0.9238059
13Pr1	K9Me3	0.9841044
13Pr1	K4Me3	0.9969566
13Lg	K27Ac	0.9824737
13Lg	K27Me3	0.9518218
13Lg	K36Me3	0.9647369
13Lg	K9Me2	0.9505661
13Lg	K9Me3	0.9927826
HPDE	K27Ac	0.9969341
HPDE	K27Me3	0.9694532
HPDE	K36Me3	0.9582686
HPDE	K9Me2	0.9820727
HPDE	K9Me3	0.9655677
HPDE	K4Me3	0.991134
38Lg_DMSO	K27Ac	0.9523149
38Lg_DMSO	K27Me3	0.9190836
38Lg_DMSO	K36Me3	0.8830686
38Lg_6AN	K27Ac	0.9515394
38Lg_6AN	K27Me3	0.9328865
38Lg_6AN	K36Me3	0.8750725
	Average	0.9331653
	Median	0.9556397

[0235] Supplementary Data 2. This provides summaries of chromatin domain calls for LOCKs, large LOCKs, and ECDs for each sample including median lengths, ranges, % genome coverage, and levels of individual histone modifications in each type of domain. Median lengths, ranges, and % genome coverage for each individual heterochromatin modification individually (irrespective of domain location) is also included.

[0236] Supplementary Data 2A: Summary of large chromatin domains detected by ChIP-seq

Samples (name, source)	Domains	Nos.	Ranges (bp)	Median lengths (bp)	Total length (bp)	% of genome	K9Me2 enrichment (ΔRPKM)	K9Me3 enrichment (ΔRPKM)	K27Me3 enrichment (ΔRPKM)
A38Per, Peritoneal	LOCKs	2,648	100,001~ 24,067,001	311,251	1,547,508,645	54.11%	0.082	0.003	0.102
	ECDs	2,021	50,499~ 8,760,499	271,999	895,772,411	31.32%	−0.116	−0.225	−0.139
	Large LOCKs	344	500,501~ 10,773,501	1,287,251	589,313,343	20.61%	0.012	0.458	−0.020
A38Lg, Lung Met	LOCKs	3,166	100,001~ 14,074,501	298,251	1,627,719,164	56.91%	0.069	−0.101	0.118
	ECDs	2,318	50,499~ 12,388,255	241,499	838,469,352	29.32%	−0.112	−0.243	−0.159
	Large LOCKs	226	505,501~ 10,428,001	1,340,751	416,070,225	14.55%	0.004	1.040	−0.152
A13Pr1, Primary 1	LOCKs	2,446	100,001~ 30,694,501	279,501	1,745,065,095	61.02%	0.038	0.063	0.124
	ECDs	2,008	50,499~ 5,944,499	255,167	914,964,363	31.99%	−0.062	−0.176	−0.221
	Large LOCKs	77	504,001~ 5,223,501	994,501	97,292,576	3.40%	−0.045	0.640	0.043
A13Pr2, Primary 2	LOCKs	2,862	100,001~ 24,749,001	298,251	1,845,941,860	64.54%	0.065	0.051	0.107
	ECDs	1,968	50,499~ 10,621,499	277,249	914,565,095	31.98%	−0.122	−0.141	−0.199
	GHDs	50	501,501~ 1,742,001	711,001	40,355,550	1.41%	0.015	0.990	−0.112
A13Lg, Lung Met	LOCKs	3,140	100,001~ 31,395,001	293,501	2,052,192,139	71.75%	0.048	0.027	0.089
	ECDs	1,935	50,499~ 3,447,581	207,499	672,028,952	23.50%	−0.125	−0.155	−0.231
	Large LOCKs	111	500,501~ 2,878,501	735,501	100,336,111	3.51%	−0.034	0.558	−0.180

[0237] Supplementary Data 2B: Summary of broad heterochromatin modifications detected by ChIP-seq

Samples (name, source)	Large histone modifications	Numbers	Ranges (bp)	Median lengths (bp)	Total length (bp)	% of genome
A38Per, Peritoneal	K9Me2	2,331	100,001~ 24,067,001	335,001	1,449,305,831	50.68%
	K9Me3 (large LOCKs)	344	500,501~ 10,773,501	1,287,251	589,313,343	20.61%
	K9me3 (LOCKs)	711	102,001~ 1,690,501	219,501	207,546,709	7.26%
	K27Me3	1,523	100,001~ 1,127,001	186,001	360,026,522	12.59%
A38Lg, Lung Met	K9Me2	2,010	100,001~ 14,074,501	255,501	979,113,009	34.23%
	K9Me3 (large LOCKs)	226	505,501~ 10,428,001	1,340,751	416,070,225	14.55%
	K9me3 (LOCKs)	189	101,501~ 1,027,001	215,000	49,777,688	1.74%
	K27Me3	2,649	100,001~ 2,197,001	244,501	882,095,148	30.84%
A13Pr1, Primary 1	K9Me2	1,696	100,001~ 1,428,501	176,501	407,098,196	14.23%
	K9Me3 (large LOCKs)	77	504,001~ 5,223,501	994,501	97,292,576	3.40%
	K9me3 (LOCKs)	1,181	100,501~ 11,051,001	322,501	573,396,678	20.05%
	K27Me3	2,211	100,001~ 30,694,501	232,001	803,104,711	28.08%

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Samples (name, source)	Large histone modifications	Numbers	Ranges (bp)	Median lengths (bp)	Total length (bp)	% of genome
A13Pr2, Primary 2	K9Me2	2,693	100,001~ 24,749,001	295,501	1,738,982,692	60.80%
	K9Me3 (large LOCKS)	50	501,501~ 1,742,001	711,001	40,355,550	1.41%
	K9me3 (LOCKS)	505	101,001~ 1,203,001	187,501	117,705,503	4.12%
	K27Me3	1,828	100,001~ 2,887,501	220,001	589,055,827	20.60%
A13Lg, Lung Met	K9Me2	3,091	100,001~ 24,749,001	283,501	1,898,353,591	66.38%
	K9Me3 (large LOCKS)	111	500,501~ 2,878,501	735,501	100,336,111	3.51%
	K9me3 (LOCKS)	711	101,501~ 1,748,001	194,501	168,847,710	5.90%
	K27Me3	2,553	100,001~ 31,395,001	237,001	1,003,794,552	35.10%

[0238] Supplementary Data 3. This provides all p-values calculated for sequencing experiments, as designated by the figure labels. Sensitivity analyses for LOCK domain calls are also included.

[0239] Supplementary File 3A: p-values for H3K9Me2 reprogramming across LOCKs

FIG. 2A: H3K9Me2 Reduction Across LOCKs (vs A38Per)				
Samples	K9Me2 Enrichment (ARPKM)	p-value (RPKM < A38Per Wilcox Test)	Mb K9Me2 Reduction (vs A38Per)	p-value (Reduced Mb vs A38Per, Chi-square Test)
A38Per, Peritoneal	0.082	NA (reference)	NA (reference)	N/A (reference)
A38Lg, Lung Metastasis	0.069	p < 2.2e-16	763 Mb (52.7%)	p < 2.2e-16
A13Pr1, Primary Tumor 1	0.038	p < 2.2e-16	1110 Mb (76.6%)	p < 2.2e-16
A13Pr2, Primary Tumor 2	0.065	p < 2.2e-16	286 Mb (19.7%)	p < 2.2e-16
A13Lg, Lung Metastasis	0.048	p < 2.2e-16	204 (14.1%)	p < 2.2e-16

[0240] Supplementary Data 3B: p-values for reprogrammed euchromatin modifications from DE genes

FIG. 2E Genes Up-regulated from Euchromatin (vs Peritoneal: A38Per)		
Sample	pvalue K27Ac (>A38Per Wilcox)	pvalue K36Me3 (>A38Per Wilcox)
A38Lg, Lung Metastasis	p < 2.2e-16	p < 2.2e-16
A13Pr1, Primary Tumor 1	p < 2.2e-16	p < 2.2e-16
A13Pr2, Primary Tumor 2	p < 2.2e-16	p < 2.2e-16
A13Lg, Lung Metastasis	p < 2.2e-16	p < 2.2e-16

FIG. 2E Genes Down-regulated from Euchromatin (vs Peritoneal: A38Per)		
Sample	pvalue K27Ac (<A38Per Wilcox)	pvalue K36Me3 (<A38Per Wilcox)
A38Lg, Lung Metastasis	p = 0.0000007224	p = 6.411e-09
A13Pr1	p < 2.2e-16	p < 2.2e-16

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A13Pr2	p = 0.0000001401	p < 2.2e-16
A13Lg	p < 2.2e-16	p < 2.2e-16

[0241] Supplementary Data 3C: p-values for reprogramming of H3K9Me3 across LOCKs

FIGS. 10A-10B: H3K9Me3 Enrichment over LOCKs		
Comparison	H3K9Me2 (Wilcox Test)	H3K9Me3 (Wilcox Test)
A38Per vs A38Lg	p < 2.2e-16	p < 2.2e-16
A13Pr1 vs A13Pr2	p = 0.1086	p < 2.2e-16
A13Pr1 vs A13Lg	p = 0.2364	p < 2.2e-16

[0242] Supplementary Data 3D: p-values for reprogramming of modifications from LOCK DE genes

FIG. 11A: DE genes Downregulated from LOCKs (vs A38Per)					
Sample	K9Me2 (>A38Per Wilcox)	K27me3 (>A38Per Wilcox)	K27ac (<A38Per Wilcox)	K36me3 (<A38Per Wilcox)	DNA Methylation (<A38Per Wilcox)
A38Lg	p = 0.99	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16
A13Pr2	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16
A13Pr1	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16
A13Lg	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16

FIG. 11B: DE genes Upregulated from LOCKs (vs A38Per)					
Sample	K9Me2 (<A38Per Wilcox)	K27me3 (<A38Per Wilcox)	K27ac (>A38Per Wilcox)	K36me3 (>A38PerWi lcox)	DNA Methylation (>A38Per Wilcox)
A38Lg	p = 0.99	p < 2.2e-16	p = 1.163e-06	p = 0.000000007195	p = 0.01699
A13Pr2	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p = 3.312e-08
A13Pr1	p = 0.51	p = 2.738e-06	p = 1.362e-06	p = 0.0004268	p = 0.99
A13Lg	p < 2.2e-16	p < 2.2e-16	p = 2.362e-13	p < 2.2e-16	p < 2.2e-16

[0243] Supplementary Data 3E: p-values for reprogramming across Large LOCK domains

FIGS. 13B-13C: Reprogramming Across Large LOCK Domains			
FIG. 13B	H3K9Me2 (>A38Lg Wilcox)	H3K9Me3 (<A38Lg Wilcox)	DNA Methylation (>A38Lg Wilcox)
A38Per Large LOCKs	p = 0.0006861	p < 2.2e-16	p = 0.000007037

FIG. 13C	H3K9Me2 (>A38Per Wilcox)	H3K9Me3 (<A38Per Wilcox)	DNA Methylation (>A38Per Wilcox)
A38Lg Large LOCKs	p = 0.00002502	p < 2.2e-16	p = 3.52e-16

[0244] Supplementary Data 3F: p-values for 6AN RNA/ChIP-seq experiments

FIG. 19A: 6AN Down-regulation of DE Genes from LOCKs with Pre-Existing Regulatory Modifications (6AN vs. other LOCK genes)				
	K27Ac (6AN > Other)	K36me3 (6AN > Other)	K27Me3 (6AN < Other)	K9Me2 (6AN < Other)
Wilcox Test p-value	p < 2.2e-16	p < 2.2e-16	p < 2.2e-16	p = 0.02238

FIG. 19B: 6AN Increase of H3K9Me2 Across Reprogrammed LOCKs (DMSO vs 6AN Over LOCKs Reprogrammed between A38Lg vs A38Per)		
	K9Me2 (6AN > DMSO)	K27Me3 (6AN > DMSO)
Wilcox Test p-value	p = 2.291e-07	p = 0.6333

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FIGS. 19C-19D: 6AN Decrease of H3K27Ac Over DE Genes Repressed From LOCKs			
FIG. 19C	K27Ac LOCK Down Genes (6AN < DMSO)	K27Ac Other LOCK Genes (6AN < DMSO)	K27Ac ECD Down Genes (6AN < DMSO)
Wilcox Test p-value	p = 0.01549	p = 0.94462	p = 0.8944
FIG. 19D	K27Me3 LOCK Down Genes (6AN < DMSO)	K27Me3 Other LOCK Genes (6AN < DMSO)	K27Me3 ECD Down Genes (6AN < DMSO)
Wilcox Test p-value	p = 0.9417	p = 0.5139	p = 0.7957

[0245] Supplementary Data 3G: LOCK sensitivity analyses

[0249] Supplementary Data 4C: Summary of DE genes between A38Per and A13Lg detected by RNA-seq and

Sensitivity Testing of LOCK Domain Calls					
Sample	Original Parameters Called Domain Lengths	Test Parameter 1 Called Domain Lengths	Overlap Analysis Percent Overlap	Test Parameter 2 Called Domain Lengths	Overlap Analysis Percent Overlap
Small Peaks Removed	5 kb	2 kb	5 Kb vs 2 Kb	10 kb	5 Kb vs 10 Kb
38Per K9me2	1449305831	1475751351	98.02%	1389543312	95.65%
38Lg K9me2	979113009	1042797151	93.44%	870263273	88.44%
38Per K27me3	360026522	383258588	92.44%	321712391	88.43%
38Lg K27me3	882095148	886275657	99.53%	863791604	97.89%
Minimal Merge Distance	20Kb	15Kb	20Kb vs 15Kb	25Kb	20Kb vs 25Kb
38Per K9me2	1449305831	1387108909	95.42%	1504614743	96.11%
38Lg K9me2	979113009	887961919	89.91%	1053584044	91.69%
38Per K27me3	360026522	314837912	86.64%	396029094	89.89%
38Lg K27me3	882095148	848539120	96.00%	906322630	97.16%

[0246] Supplementary Data 4. This file lists all differentially expressed (DE) genes detected in each sample by RNA-seq, including level of expression, p-values, directional changes, and chromatin domains that each DE gene mapped to. Analysis of recurrent DE genes detected across distant metastatic samples and between control (DMSO) and 6AN treated cells is also reported.

[0247] Supplementary Data 4A: Summary of DE genes between A38Per and A13Pr1 detected by RNA-seq and mapped to chromatin domains (data not shown—publicly available on the World Wide Web at nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true #supplementary-information by clicking link “Supplementary Table 7”).

[0248] Supplementary Data 4B: Summary of DE genes between A38Per and A13Pr2 detected by RNA-seq and mapped to chromatin domains (data not shown—publicly available on the World Wide Web at nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true #supplementary-information by clicking link “Supplementary Table 7”).

mapped to chromatin domains (data not shown—publicly available on the World Wide Web at nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true #supplementary-information by clicking link “Supplementary Table 7”).

[0250] Supplementary Data 4D: Summary of DE genes between A38Per and A38Lg detected by RNA-seq and mapped to chromatin domains (data not shown—publicly available on the World Wide Web at nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true #supplementary-information by clicking link “Supplementary Table 7”).

[0251] Supplementary Data 4E: Summary of DE genes between A38Per and A38Lv detected by RNA-seq (ChIP-seq not performed for chromatin domains) (data not shown—publicly available on the World Wide Web at nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true #supplementary-information by clicking link “Supplementary Table 7”).

[0252] Supplementary Data 4F: Summary of DE genes recurrently up/down-regulated across primary tumor precursor (A13Pr1/2) and distant metastatic subclones (A13Lg,

A38Lg, A38Lg) vs. A38Per (data not shown—publicly available on the World Wide Web at [nature. com/ng/journal/ v49/n3/full/ng.3753.html?foxtrotcallback=true](https://www.nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true) #supple-
mentary-information by clicking link “Supplementary Table 7”).

[0253] Supplementary Data 4G: Summary of DE genes between control (DMSO) and 6AN treated A38Lg cells detected by RNA-seq and mapped to chromatin domains

(data not shown—publicly available on the World Wide Web at [nature.com/ng/journal/v49/n3/full/ng.3753. html?foxtrotcallback=true](https://www.nature.com/ng/journal/v49/n3/full/ng.3753.html?foxtrotcallback=true) #supplementary-information by clicking link “Supplementary Table 7”).

[0254] Supplementary Data 4H: Comparison of overlaps between DE genes between matched lung and peritoneal subclones (A38Lg vs. A38Per) and DE genes between control and 6AN treated A38Lg cells.

38Lg up/ 6AN down Coregulated/ 6AN down: 1032/1968 (52%) Coregulated/ 38Lg up: 1032/4368 (24%) (Coregulated: genes present in both 38Lg and matched 6AN datasets)	Genes	38Lg down/ 6AN up Coregulated/ 6AN up: 915/2192 (42%) Coregulated/ 38Lg down: 915/4402 (21%)	Genes	38Lg up/ 6AN up Coregulated/ 6AN up: 379/2192 (17%) Coregulated/ 38Lg up 379/4368 (0.09%)	Genes	38Lg down/ 6AN down Coregulated/ 6AN down: 277/1968 (14%) Coregulated/ 38Lg down 277/4402 (0.06%)	Genes
	GDF5OS		CCT6B		WDR25		RP11- 400K9.4.1
	EDN2		JAK2		TOMM7		C14orf105
	C1QTNF2		TMEM63A		IGFBP1		CCK
	CLDN2		FAM101A		SLC9A3R2		RP11- 255B23.3.1
	FAT2		KRCC1		C7orf60		CFTR
	RBM24		ERRFI1		ZFYVE28		CD22
	KMO		LRRC6		CTSD		RGS7
	TPRG1-AS1		TNFAIP2		GATA6		ERVMER34-1
	DOCK2		MAST3		RP11- 480A16.1.1		CCNI2
	FAM101B		ABCC10		GXYLT2		ADAMTS14
	KRT7		RP11- 65J3.1.1		AC002472.8.1		ANO1
	TCF7		SLC12A7		C3orf23		PADI3
	RP11- 845M18.3.1		WASH7P		TMEM43		CHST4
	C4orf49		ANKRD13D		FUT11		FER1L6
	SLC1A3		ZBTB7A		IFFO1		TNNT2
	DMC1		WASH6P		SLC1A4		RP11- 597D13.9.1
	TMSB15A		IL32		DDT		RP11- 6F2.4.1
	LHX4		RNF217		IDI1		MYO5B
	KIAA1324		EFNB1		SARM1		KRT23
	ITPRIPL1		KHNYN		HOTAIR		MUC4
	PPAPDC1A		TMEM59		TSC22D1		RP11- 346D6.6.1
	RGS5		GAS8		NBAS		ITGAM
	RP11- 618G20.2.1		ABC7- 42389800N19.1.1		ENG		PTPRZ1
	RIPPLY1		RASSF7		SSTR5		GPR116
	UHRF1		MRPS6		CRIP2		CYP24A1
	SUSD5		RP11- 353B9.1.1		NUDT18		DHRS9
	KRT80		GRAMD4		RPL31		GRPR
	KRT32		SERINC5		KIAA1143		AIF1L
	SLC26A7		TMEM102		DNAJC3-AS1		CGN
	TFPI2		TMPRSS5		C1R		RP11- 314P12.3.1
	RP11- 314P12.2.1		TMEM63B		GAB2		CYFIP2
	MMP7		FAM109A		FAM174B		ATP6V0A4
	KLHL23		TLR4		SYT11		MPP7
	C1orf110		C20orf96		DET1		KREMEN1
	GRIN2A		SMAD6		TBC1D8B		TTN
	PADI2		VAMP4		TAZ		LEF1
	AL162759.1.1		MYO1E		UCP2		SP140
	MYBL2		DVL1		RP4- 647C14.2.1		SP6

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GLYATL2	AC069513.3.1	AARS	ANXA10
RRM2	ITPKC	DALRD3	NYAP2
E2F2	USP18	SEMA3C	MCOLN3
CPA4	RAB33B	ATP2B1	DENND2A
HHIP	RP11-325F22.3.1	THTPA	HSH2D
MYPN	ADHFE1	GPER	C1orf106
RP11-150O12.6.1	LENG8	ITPK1	EVL
CLSPN	RILPL2	IL1RAP	C7orf58
WDR69	TSNARE1	RPL29	RP1-95L4.4.1
RP11-150O12.1.1	SPIRE2	EFHC1	IL11
AKAP12	MMAA	CRTAP	DDN
DOCK10	SLC15A3	B3GAT1	snoU13
PSG5	PLEKHM1	SEL1L	SLC27A2
SPC25	RP11-403I13.8.1	ME1	DGAT2
TM4SF4	STARD10	TBC1D9B	SHANK2
FAM71D	SNPH	EEF1A1P5	HNRPCP
OLFML2A	EME2	VKORC1	AC005083.1.1
OXCT1	DHRS3	SNX21	S1PR3
FAM111B	MX1	KDELC1	ARL14
snoU13	RRN3P1	AIP	SAMD5
AC073130.1.1	FAM100A	RP1-34B21.6.1	ARHGDIB
CCNE2	DDO	OSBPL6	GNG4
CDC25A	TPRN	C4orf34	CDC42BPG
RP3-324O17.4.1	CD58	FAM18B2	SMO
MCM10	PLCG2	TXNDC15	TIAM1
RP11-424C20.2.1	RP11-220I1.1.1	LIN7A	NMU
MSRB3	ZNF554	CTD-2287O16.1.1	SOX6
HEATR7B1	GLI4	FLYWCH1	PPFIBP2
TP73	TRAF1	C11orf2	LY6D
PRKDC	ANKRA2	CHST12	PLAC1
DLEU2	SYNGR2	FTX	SYTL5
CTB-164N12.1.1	FAM113B	CCNO	SELL
SDPR	CROCC	AL844908.5.1	MYO5C
POLE2	HIVEP2	RP11-996F15.2.1	UNC5A
MEST	RHBDD2	CPEB2	SPNS2
ASTN2	GATS.1	PDIA6	BIK
LAMP5	RALGPS1	STON1	PPYR1
MKI67	VWA1	ARRDC3	FAM46B
TMEFF2	HEATR7A	C7orf23	COL17A1
CPA5	HLA-K	TESK1	MACC1
ANKRD18DP	MAF1	HEG1	DNAH12
ESCO2	TMEM8A	RP11-85K15.2.1	CYP2J2
TUBA1B	DDX58	MTHFD2	LDLRAD3
WNT7A	PYROXD1	RWDD2A	RAB6B
MCM4	CYP27C1	RPL10	BTBD11
AKAP5	RNF145	PHACTR1	PLA2G7
TYMS	ZDHHC14	NUDT22	TBC1D30
CDCA2	KLF7	TRIM4	SNX25P1
CAP2	KYNU	RHOG	TESC
RP11-527N22.2.1	LDLR	EIF1B	COL1A1
RP11-297M9.2.1	ZDHHC11B	SSR4	FERMT1
DEPDC1B	AC007383.3.1	ARG2	MED12L
DPF1	SRCRB4D	TCEAL1	RP11-582J16.5.1
RP11-554I8.2.1	LRP5L	ZFPL1	BRI3BP
RP11-33N16.1.1	FOXC1	RGS10	LAD1

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MAP3K14	AACS	MEIS3	C16orf74
EXO1	SLC44A2	LLGL1	CD163L1
RP11-253E3.3.1	SEPP1	C9orf37	ANXA8L1
FABP6	ARHGAP25	USE1	AEN
RP11-184I16.2.1	STAT6	YPEL1	C8orf46
FOSL1	RP11-27I1.2.1	PLA2G15	CDH1
CCDC99	VPS28	RPS27	XDH
AC016831.7.1	R3HDM2	VEGFB	XK
FJX1	C11orf35	CELSR3	ANXA8L2
SCN5A	ARHGEF10L	EPHX1	COL12A1
RP11-687M24.3.1	PSMD9	TINF2	PPARGC1B
ILDR2	METTL7A	ZNF70	C6orf132
MUC5AC	TMTC2	C3orf18	TTC3P1
AC092614.2.1	NFE2L1	PITPNC1	GALNT6
APOBEC3B	C5orf13	ISL1	SEMA7A
RP11-54A9.1.1	ZNF768	CORO6	F2RL1
NCAPG	SIK1	NEK3	ALDH1A1
ATAD2	OSCP1	RP11-313D6.4.1	CRABP2
SMC4	ACADS	RPL27A	CA2
FUT9	HERC6	RP11-216F19.1.1	FGFBP1
ZNF488	CLDN9	ABCA3	SNAI2
TK1	RHEBL1	PITPNM2	PLS1
CTD-2023N9.1.1	CHST6	LGR5	KLHL13
HELLS	VMAC	AGA	RP5-862P8.2.1
MMP24	AQP3	FAM113A	RAB38
SLITRK3	DNAL4	GPR37	SLC7A8
ELOVL2	HBP1	VASH1	WNT3
SERINC2	ZG16B	CAMKK1	SARDH
AC002066.1.1	FAM100B	MAST1	TPM1
DIAPH3	EREG	RP11-574K11.20.1	RASEF
RP11-291L15.2.1	ZNF862	RPL12	SKAP1
GIN51	NARFL	NSUN5P2	TRIM14
RP5-968P14.2.1	SWSAP1	RP11-477I4.3.1	CTSL2
DHFR	YPEL5	NSUN5P1	GK
ROR1	SLFN5	HAUS4	VEPH1
MCM3	CCNDBP1	TMEM234	RP11-800A3.4.1
RP1-140K8.5.1	PAQR8	MRC2	FGFR3
GLYATL1	SLC25A27	PYCR1	PADI1
CDCA7	GOLGA8B	SLC26A6	CDS1
CSMD2	IKZF2	C3orf78	JPH1
RBPM52	HSD17B11	CD68	FAM81A
RBL1	LMBR1L	AP003068.23.1	GALNT12
RP11-677M14.3.1	KCNJ14	PAK3	DPY19L2
CCBE1	DDR1	GMPPA	ITGBL1
CCND1	C7orf53	EEF1A1P6	SYNPO2
E2F8	RELB	AC093673.5.1	RP11-157P1.4.1
CDC45	WASH3P	CBS	ANKRD22
LMNB1	PDCD4	MIF4GD	GRHL1
C11orf41	NR1D1	CNPY4	KRT15
BEST3	NCOA7	GGT7	RP11-93L9.1.1
CTPS	GOLGA2	LRP1B	C15orf62
RASSF10	CEBPG	FAIM2	AADAC
POLA1	C2orf15	ZNF333	AC112229.1.1
TUBB6	KLF13	AGXT2L2	TRIM59
HOMER1	GEMIN8	B9D1	DNAH10
CD200	ARRDC2	EEF1A1	CTD-2021J15.2.1

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CENPM	SLC39A11	ATRNL1	KRT19
ERCC6L	SAT2	PAOX	KIAA1199
HMMR	ZER1	CTSA	GPM6A
CENPW	KIAA1407	SLC45A1	AKAP6
RP11-298I3.4.1	OAS1	RAB3IL1	GPR110
RP11-259P6.1.1	AC004410.1	C6orf1	QPCT
THBS1	DGAT1	CTD-2314G24.2.1	RP11-382A18.1.1
GPR63	RNF103	CYP4X1	METTTL7B
DTL	EFHC2	HCFC1R1	PHACTR2
RP11-101K10.8.1	C17orf69	FAM70B	PLEKHA7
RP11-204J18.3.1	LINC00493	SHMT2	PLAU
WDR4	ST6GALNAC2	SEC14L5	SLC4A3
BRIP1	KLHL31	CLK4	KRT18
LMNB2	LGMN	UGDH-AS1	C3orf52
C14orf49	LLGL2	ITGA11	SYTL2
EIF6	ENDOV	CGREF1	PKDCC
ARHGEF26	CST1	RPL22L1	ASB9
ETNK2	RP13-516M14.1.1	FKBP2	SLC22A20
POLQ	LIPA	OXA1L	IGSF3
RP11-573I11.2.1	FAM13A	C10orf102	CLDN10
APLN	ST3GAL1	DNASE1L1	MBOAT1
RP11-304F15.3.1	APOBEC3G	HIGD2A	TMEM169
NDC80	PLAUR	MLLT3	SYK
MCM6	ACYP2	AP001496.1	MAST4
CAV1	RTN2	FAM156B	SDC1
RP11-53M11.3.1	HIF1A	TRPT1	RP11-613M10.6.1
MARK1	DNASE2	RP11-571M6.6.1	PPM1H
CTA-445C9.15.1	CA11	RP11-243J18.3.1	GCHFR
USP13	SLC7A7	SGSH	RHOD
RP5-1172A22.1.1	ABCA10	CTD-3074O7.5.1	EZR
UCN2	PTK6	CNTNAP1	RP11-303E16.2.1
MCAM	RPS6KA2	TNFRSF10C	RCC2
ZWINT	WHAMMP3	B3GNT1	KDR
LINC00460	ENGASE	LRP1	RP11-416I2.1.1
BIRC5	LPCAT4	P2RX6	GPR65
FLNC	EPS8L3	SSBP2	BCL7A
SYNE2	TMEM198B	ZC3H6	PPCDC
HMGN2	PRICKLE3	CCNG1	NBPF10
ADCY3	PDE4DIP	TBC1D4	AC108463.1
PDSS1	GPR108	MAGED2	CADM4
CDCA3	TMC4	DNAJC1	SPTBN5
AXL	PTP4A3	PCK2	CUEDC1
KIAA0101	ITGAX	MANF	CMTM4
NAV3	TSSC4	ZCWPW2	CDK5R1
KIF11	ANO9	C1orf213	C19orf21
LPCAT2	PPP1R3B	CACNG6	C1orf116
KIF4A	ANTXR2	AC004540.5.1	UNC5C
CTD-2334D19.1.1	ARHGEF3	TSLP	ALDH5A1
EML5	C10orf32	RP5-1103G7.4.1	MB21D2
NME1	PGM2L1	C1S	RP11-7K24.3.1
ARHGAP11A	HLA-H	SPATA25	DSG2
MCM2	TUBG2	PPAP2A	PLD6
SLC5A11	SLC16A3	RP11-390P2.4.1	GALNT5
CHN1	CCNL2	RP11-539L10.3.1	ZNF185

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KIF23	SGK1	SH2B2	P2RY2
ARHGAP19	MEF2C	NEURL2	PLCXD1
SLC8A1	CAMTA2	KLHDC2	AIG1
FBN2	HLA-F-AS1	SHISA2	PALM
FKBP5	PIWIL4	PXK	PKP4
IGSF9B	LSS	WBSCR27	ZNF462
KIF15	PCMTD2	DSEL	SAA1
NUP210	CFI	TRIM46	SPATA13
AS3MT	ARL4D	CTSF	LAMA3
TNFAIP8L3	MVK	KLHL35	NID1
PM20D2	CDH17	PDIA5	PEAR1
OPN3	MAFF	PLK3	PYGL
RRM1	CACNB1	ZNF815	USP40
DNA2	C2orf63	NICN1	DIS3L
DEPDC1	IL15	SCAND1	ID2
PDE1A	ST3GAL6	CALHM3	FUT1
ALYREF	C9orf7	RP5-827C21.4.1	PRSS3
MORC4	ENDOU	LETMD1	NMNAT2
FBXO5	PIM3	JAKMIP3	DNAJA4
DPY19L2P1	CLDN7	LZTFL1	DOCK9
CENPF	DAB2IP	CTNS	DDI2
GPR19	TARSL2	RP11-280F2.2.1	PLAC8
SYCE2	REC8	PDK1	FBP1
GIN54	FAM160A2	MBL1P	RAG1
XRCC2	ZFP36	LHPP	PRPF4
DARS2	RP11-73K9.2.1	BLVRB	RPS6KA1
HMGB3	STARD4	CALHM2	DSP
KIAA1524	SGK2	RP11-277P12.20.1	SLC22A5
NUP188	CIR1	PTCHD2	AADAT
DPF3	MAPK8IP3	UST	FARP2
STMN1	USP20	IRF2BPL	MCC
WDHD1	RP11-149I23.3.1	PLK1S1	ZNF658
NCAPD3	RP11-108M9.4.1	TTC39B	B3GNT5
CCNE1	HIP1R	ABHD14B	NT5DC3
TRPV2	CYP4F3	DLG4	ABCC9
CDKN3	SH3YL1	PCDHGA7	PNPLA4
PRR11	TNFSF12	MAGED1	ZNF717
CSRP2	TMEM8B	LEPREL2	RAVER2
COTL1	IGSF8	LOXL2	MREG
MSH2	PTK2B	SLC43A2	RBP1
KIF18B	NR4A2	ZEB1-AS1	RAPGEF3
SKP2	SPIRE1	LENG8-AS1	CNKSRR3
ADORA2B	DLL1	C17orf72	RP11-757F18.5.1
GFRA3	TMED1	C16orf93	PRSS12
ZNF724P	SQLE	ARHGAP4	HOOK1
RASGRF1	ABCC6	CNTD1	COL16A1
FAM83D	VSIG10L	PCDHGA3	MST4.1
BUB1	FA2H	STK32A	CMIP
MTHFD1	OSBPL7	SLC29A4	SMC5
DHX9	KCNIP3	LEPRE1	ANXA2P2
NRG1	LYNX1	HOMER3	STEAP2
PCNA	ULK3	MCEE	NUDT14
UTP20	HINT3	XBP1	MMD
MIR17HG	FDFT1	ALDH1L2	ANKRD56
ANLN	TRIB1	MAP2	SURF2
RUVBL1	NPDC1	NNMT	SH2D3A
MYO1B	VEGFC	NUCB1	MAP3K1
NCAPG2	GPRC5C	KLRC2	UACA
FH	SIGIRR	TMEM158	TAF2
GNG11	TCEANC	SEPT7L	DCBLD2
GS1-465N13.1.1	MKNK2	RP11-161M6.4.1	RP11-295M3.1.1
MT1L	N4BP2L1	AC079922.3.1	NAALADL2
TNS1	KCNMB4	PCDHB7	CNTRL
BARD1	RP5-1182A14.3.1	HSD17B7P2	CD82

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THOP1	GET4	NAT6	FBXL19-AS1
SKA1	FKBP10	PTX3	RHBDL2
C1QL1	RDH10	ZNF836	RP11-448G15.3.1
C11orf82	SLC22A15	TMEM106A	RP11-357H14.19.1
CCNA2	CCDC57	TFF2	TUBGCP5
SLC35F3	SHB	CYP1A1	INADL
C10orf140	TMPRSS3	PPP1R3E	C9orf64
NEXN	NYNRIN	TMEM120A	ARAP2
ASF1B	TNFRSF9	POLN	KRT16
PRTFDC1	SESN3	RAPGEF4	ALS2CL
FAT3	A2LD1	RABAC1	
H2AFX	RP11-475N22.4.1	CACNA1G	
PRIM1	ZNF628	LOX	
SNX18P3	JAG2	FAM151B	
RP11-973F15.1.1	RP11-244H3.1.1	PTPRM	
ADRB2	GOLGA8A	CLEC2B	
FBXL13	TP53INP2	KDELR3	
CDC20	WASH4P	ISYNA1	
CENPK	TNIP1	RASL11A	
CDCA4	RP4-659J6.2.1	CALR	
ATAD5	CCDC126	RP11-108K14.4.1	
SMC1A	PTPRB	METTL12	
FAM196B	U6	BNIP3	
UGT1A1	SPRY3	FAM175A	
CTD-2574D22.2.1	MMP15	CRLF2	
MCM5	SLC23A3	TMEM231	
MELK	TBC1D3F	CRELD2	
LYPD1	ABCA5	PLXNA3	
DLGAP5	RP4-798A10.2.1	AC026202.3.1	
NUSAP1	CLDN15	ASNS	
TUBBP1	TMC7	RASIP1	
MYH10	TCTEX1D2	SFRP5	
MIR155HG	NOS3	RP11-66N24.3.1	
CCDC138	ARHGEF16	ABCC3	
ERCC2	TPBG	RIBC1	
KIF20A	FAAH	HSP90B1	
TMEM14B	TMEM150A	HYOU1	
H2AFY2	LRRC56	BAMBI	
OXTR	CEP85L	TPP1	
RDM1	MIA	ZCCHC24	
PLEK2	CDH6	FAM161B	
SHCBP1	PON3	AC004080.12.1	
GJC1	CCDC92	MAPT	
SNRNP25	PTPRH	AC018755.11.1	
KCNQ5	FOXQ1	PLOD2	
ODC1	J01415.23	MAN1A1	
EBNA1BP2	PPFIA3	NUCB2	
FABP3	SMPDL3A	DERL3	
MCM7	AC007283.5.1	SCN1B	
CHAC2	ALPPL2	FN1	
BORA	MTHFR	PLCD1	
ASRGL1	WASH2P	CCDC85B	
VIPR1	RSAD2	BTBD19	
PTTG1	LIMA1	AP000769.1	
ACTB	PPP1R16A	AC147651.3.1	
TOP2A	PIK3C2B	ATHL1	
NEIL3	XAF1	CYP2E1	
ALDH1B1	KIAA0513	FAM182B	
CEP250	GRAMD1C	PDIA4	
FAM131B	GLDN	ZFP2	
FAH	GPCPD1	SLC25A29	
CIT	CABP4	RAB24	
CDCA8	NXNL2	RP11-755F10.1.1	

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CDK2	SEMA4C	QPCTL
EZH2	PHYHIP	ALDOC
PSMC3	AC093734.11.1	AKR1C1
FGGY	IRF9	LRRC29
NXPH4	ZNF517	RP11-
		307O13.1.1
FRMD4A	RP5-	SEMA3F
	1187M17.10.1	
CSPG4	HIST1H1C	RUSC1-AS1
CSE1L	ANKRD42	EVI2B
KIF20B	ZC3H12A	TIE1
C2CD3	JUND	HSPA5
PPIAP29	SEMA4B	TSPYL2
CTA-	CTAGE5	HERPUD1
221G9.10.1		
TPX2	NLRP1	AQP2
DKK1	GNE	RP4-
		794H19.2.1
PFAS	LTB4R2	C2orf16
PBK	BBS12	AKR1C2
TRIP13	DICER1-AS	ANGPTL4
CCDC18	CMPK2	HCN2
UBE2T	CALB2	NOG
AL357673.1	DHRS2	AKR1B1
PDE12	LRFN3	ST7-AS1
NUP155	PROC	EVI2A
ARNTL2	PROS1	AC022007.5.1
POLD2	TMEM80	ZNF575
NCAPD2	RALGDS	RCN3
VCL	SLC6A8	PCDHB15
CCDC85C	TMC6	RP13-
		895J2.7.1
FANCD2	HMGCR	C9orf150
RP11-	TBC1D17	ANGPT1
117P22.1.1		
MYLK	TBC1D3	ARSA
RP11-	GRN	RP11-
394B2.4.1		49I11.1.1
PRLR	ROM1	PRPH
AP2B1	MID1IP1	AC004383.5.1
RASSF2	C15orf61	P4HA1
PRKAG2	SPATA20	ANGPTL2
COL4A6	FAM78A	AC002480.4.1
PSMC3IP	STAT2	MTMR9LP
ARRB2	PYGM	CRELD1
CAMK4	RHBDF1	UPB1
TSPAN2	LRCH4	AC002480.3.1
RFC2	PODN	RP11-
		554A11.9.1
PLK2	SPRY4	GALNT9
SRRT	CITED4	AL137145.2
WNT7B	LIPG	VLDLR
ADORA1	CAPS	IFITM10
C22orf29	EIF1	LDHD
HTR1B	CYP4F12	ERO1LB
ITGB8	AKAP17A	NTNG2
ZNF660	ACTR1B	NPY1R
EPB41L2	WDR66	
COQ3	AP001468.1	
AASS	AP001372.2.1	
PAICS	FAM214A	
CENPI	GSN	
CASC5	FAM193B	
TUBA1C	TPRG1L	
NUDCD1	ARHGEF2	
L2HGDH	TNFAIP3	
CHML	CDA	
CCNB1	PLA2R1	
RP11-	ZSWIM4	
799O21.1.1		
DNMT1	IRAK2	
CDK1	LYZ	
RP11-	TJP3	
673C5.1.1		

-continued

NUP37	C8orf55
TPMT	CTB-
	131B5.5.1
AURKA	AC103810.1
PEA15	PCDHGB2
NRGN	KLC4
PLK1	RP4-
	541C22.5.1
ZNF124	SIX5
CGNL1	SEL1L3
SNRPD1	MIR29C
PTPN14	MZF1
CDC7	ADAM8
DERA	WDR45
PYGO1	ZNFX1-AS1
LRRCC1	AC017099.3.1
FAM173B	ARSD
AHCY	DLX4
UBAC2-AS1	HK2
HIGD1A	HS3ST1
GMNN	ABCG1
NCAPH	CCDC146
CEP128	EGLN3
SLC38A5	CLK1
TUBB4B	UPK3B
RP11-	SLC16A6
512F24.1.1	
BICC1	HLA-F
HMGB1P5	NT5M
NUF2	BTG1
VCAN	C7orf63
FAM64A	RASSF9
DNAJC9	CTSL1
RANBP1	ENDOD1
C4BPB	FAM116B
C9orf140	KNDC1
SKA2	PPP1R3F
RP11-	LARP6
181C3.2.1	
PKMYT1	FBXO6
RFWD3	CCDC69
TCOF1	PRRT1
KNTC1	CTD-
	2258A20.4.1
DKC1	TSPAN1
MPP2	RP11-
	362F19.1.1
CCNF	SLC2A10
C1orf112	CTD-
	2341M24.1.1
AC046143.7.1	C2orf81
STMN3	FZD4
OIP5	RAB40C
DGKH	EXD3
NRM	IFIT3
RFC3	PCDHGC5
RBBP8	RP11-
	285F7.2.1
HTR7P1	RHBDF2
NMT2	PRICKLE4
ARHGAP11B	RABL2A
DPYSL3	WBP1
TCAM1.1	LAMB3
YBX1	RP4-
	697K14.7.1
HSP90AA1	TMEM91
ACTA2	PCDHAC1
MPP5	TRIM2
C8orf84	C16orf7
ASPM	RTP4
LRRC8C	KIAA1875
TBC1D7	RP11-
	540D14.6.1

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RP11-462L8.1.1	JUP
ZNF367	NEU1
RP11-956J14.1.1	DDX60
CCT5	HOOK2
C6orf52	BCO2
NEK6	TBC1D8
TSPY26P	PRRX1
MRPL1	KRTCAP3
FARSB	PINK1
PKP2	NFIL3
CAND2	LCN2
MRT04	TYMP
KIAA0586	RP11-429J17.2.1
DEK	WARS
FAM54A	COL6A1
FEN1	LZTS2
RP3-510D11.2.1	NPIPL2
SMC2	LCA5L
RELT	KB-1460A1.5.1
SGOL1	CYP4V2
ANKRD1	FAM59A
SUV39H2	SEZ6L2
KDELC2	PTPRE
HMGB2	NR2F6
ATRIP	COL11A2
SACS	RENBP
DEPDC7	AQP6
HMGB1	ULBP1
RP11-380J14.1.1	CX3CL1
POP1	CCDC149
GFAP	ASAH1
NUP205	TNFRSF14
CENPA	C12orf63
TPGS2	C12orf57
CD109	PSPN
WDR3	RP11-263K19.6.1
NOP56	MAPRE3
PTRF	RP11-44N21.1.1
PHIP	TNFSF13B
QSER1	FMO5
FAM86A	PRX
POC1A	GSDMB
CYB5RL	JUNB
H2AFZ	HMGCS1
AC092329.1	CSF2RA
RAD51AP1	HEXDC
CCP110	PLXND1
CEP55	NFAM1
ZNF347	PER1
NCS1	CALCOCO1
RAD54L	DRAM1
MDM2	AOC2
SMTN	GAS7
CDC25C	IFIT2
TFDP1	MTRNR2L9
NEURL1B	TSTD1
TBCD	NR4A1
KRBOX1	TM6SF1
SAE1	CBLB
CHEK1	CCDC24
RP11-678B3.2.1	FAM66C

-continued

REV3L	C9orf16
SPC24	KDM6B
TMPO	TBX6
DHRS4L2	MMP28
MYOF	ACSL1
TUBA4A	G0S2
ADAT2	CDC42EP2
RP11-348A11.4.1	SCD
BST1	NFKBIA
FOXM1	PLEKHF1
AC027612.6.1	AGFG2
RP11-1334A24.4.1	LEPREL4
UBE2N	ULK1
PRIM2	ADAMTS13
S100A2	CAMK2N1
NEGR1	PDE7B
HNRNPAB	AC073343.1
CCDC41	AC002117.1.1
CAMK1	SPSB3
GTSE1	SLC5A12
MMP2	PLA2G6
BOLA3	HOXD4
SYT15	HS3ST3B1
BCL2	JAK3
ECHDC3	C10orf10
KIFC1	NFKBIZ
G3BP1	DUSP8
PFN1	NEIL1
RAET1K	ECEL1P2
MNS1	RP11-783K16.13.1
SIRPA	KLHDC1
RP11-152N13.12.1	MST1
PORCN	ROBO4
NOC3L	SYT17
RAD18	NR1H3
ESPL1	GDPD3
PTER	CEBPB
RP11-1277A3.2.1	SDCBP2
NUP107	NOV
SGOL2	DNAH7
DIO2	IER5L
ZNF726	FANK1
SFXN2	ORAI3
FRMD6	SLC5A3
CENPN	B3GALT4
NT5DC2	ZDHHC1
RFC4	KIF27
NOP16	GIMAP2
TUBG1	CDC42EP5
DZIP1	AC006028.9.1
IPO5	PRSS16
LHFP	TGFBR3
ARHGAP22	FASN
TMEM48	ATP1B1
RPA3	C17orf108
TMCO7	EPHB6
THSD1	BCL6
AC013461.1.1	RP4-811H24.6.1
CCDC165	RP11-496I9.1.1
MAGOHB	CXCL2
EFEMP1	CACNA2D4
EED	PPM1K
KIF14	C17orf103
NUP35	MXD1
DTYMK	MAGIX
SSX2IP	PTPRCAP
CELF2	DPM3
NETO2	EVPL
PHLPP2	SLC2A13

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PTBP1	IL1B
FKBP3	KCNE4
URB2	INPP5J
EEF1E1	FRAT1
BEGAIN	DUOX2
CACYBP	FBXL15
HMGN5	C15orf48
HJURP	FAM86FP
DBF4	NR3C2
XRCC3	STX1B
TMCC3	RP11-
	273G15.2.1
SPAG5	RASSF4
AC108488.3.1	P2RX4
ATOX1	TPD52L1
SLC25A3	RP11-
	420G6.4.1
TUBB	P2RY11
CCDC152	ADCK3
CENPL	PARP10
GAS6	AC103810.2
RP11-	HERC5
540A21.2.1	
NXT2	TFEB
LRR1	FAM84B
E2F1	AC093627.10.1
RP11-	OAS2
122A3.2.1	
KCNH4	ARHGAP9
MYEOV	FBXO24
TMEM194B	CD34
RANGAP1	CD14
CNTF	OTUD1
NASP	ICA1
MAP7D3	AC005152.2.1
RP1-	WDFY3-AS2
239B22.1.1	
DUS2L	HIST1H2AC
FKBP1A	SELPLG
NOLC1	PDGFRB
ZNF702P	RP11-
	566K11.1.1
CDCA5	SHC2
IMMP2L	CES3
SUPT16H	GBP5
MIR621	MYL5
RP11-	JHDM1D
64D22.2.1	
CEP97	YPEL2
ITGB3BP	DHX58
ERCC8	IFIT1
WDR62	RP3-
	395M20.8.1
ZNF239	DFNB31
RP11-	LYG1
680F8.1.1	
GNL3L	FAM66D
MAPKAPK3	RP5-
	882C2.2.1
ADK	IL1R1
CENPV	LTB4R
UTRN	PDXDC2P
PSMD1	IL4I1
GTF2H3	FOXO4
USP49	AC127496.1
C3orf26	ANKZF1
CYP26B1	CSF2
JAM3	LENG9
SPA17	AC097500.2.1
BTG3	CCDC19
MRPL15	FURIN
KPNA2	IFIH1
RRP15	GBP4
HSPB11	CCNG2
C17orf89	FZD1
HAT1	SNHG5

-continued

UBE2S	PDZD7
RBM12B	DAPK2
MRPL47	C20orf195
MSI2	C9orf163
KIAA0020	PCSK4
PTPN1	KIF26B
GLRX3	SERPINE2
CFL1	MSMO1
CDH24	CTC-
	523E23.1.1
HNRNPA3	HIST1H3E
XYLB	PAQR6
DRAP1	P4HA2
ACTR3B	AOC3
CHCHD3	IRS2
H2AFV	MAFB
NSMCE1	MXD4
COLQ	DNER
CEP41	MDGA1
NUTF2	CH25H
SMARCC1	TST
KIF2C	RARRES3
ATP5G1	UNC13A
ISPD	MST1P9
CHTF8	DNAH2
IPO9	KCNK5
AC026271.4.1	PPIL6
TTF2	RP11-
	1391J7.1.1
DHRS4	RHPN1
AC068282.3.1	FOSB
SF3B3	RP11-
	202P11.1.1
GSG2	MIR29B2
AHCTF1	CEACAM1
LPHN3	RNF24
KIF22	YPEL3
IMMP1L	GPT
SNRPB	CALML6
DCLRE1A	RP1-163M9.6.1
ZNF681	NPPA-AS1
CDH2	PELI2
GPR125	FAM71E1
ANKRD18A	BPI
RP11-	RASSF5
110I1.12.1	
SCD5	TMEM53
ARMC10	ABTB1
AC009948.5.1	UCN
KIF5C	IDUA
RP11-	GDPD1
381E24.1.1	
PACSIN3	RP4-
	758J18.10.1
GNB4	BMF
RP11-	TRIB2
58E21.3.1	
PARP1	DYRK1B
HSPA4L	SCNN1D
PRPS1	CLIP2
PPIL1	INH1A
XRCC6	FAM47E
COQ2	LRRC24
CBX2	GFI1
LBH	EPB41L4A-AS1
RNASEH1	MVD
RFT1	THBS3
C1orf114	CARNS1
ITSN1	C16orf79
ATXN10	SLC16A13
WHSC1	CTD-
	2292P10.4.1
RP5-	PDE5A
991G20.4.1	
SAAL1	CTC-
	378H22.2.1

-continued

FER	MUC1
RP1-152L7.5.1	AC008440.10.1
MCM8	PCSK9
C4orf10	CTD-2547L24.3.1
CROT	ICAM5
GTF2H2	MUC20
BAG2	PPP1R3C
METTTL8	KLF4
SEC14L2	NINJ1
C20orf94	CLIP3
HDAC8	CTD-2517M22.14.1
PRMT3	AC022098.1
HAUS1	ABCA6
POLR3G	BBC3
CCT2	RP13-15E13.1.1
RP11-14N7.2.1	CXCL3
AC034193.5.1	RP11-369J21.5.1
CNTLN	PRR15
CCDC34	C14orf45
DLX1	TMEM198
ABCC4	ZNF425
RFC5	GAL3ST1
PSMC5	SLC1A7
CSTF2	ISG15
LIG1	ATF3
SLC19A1	ICOSLG
BRCC3	ATP8A1
ZW10	LINC00324
AFAP1L1	FBXO32
CDC123	EXOC3L4
RP11-521B24.3.1	SRCIN1
LRRC58	HSF4
ALMS1	DBP
FAM111A	CDH3
PUS7	AC021593.1
DYNC1H1	RNF152
EFNB2	ISG20
BCAS4	KIAA1683
MYH9	RP5-885L7.10.1
HSPA14	CARD14
TMEM56	PIP5K1B
ZDHHC2	NCF2
KIF18A	PPP1R32
CDK4	RAB17
CKAP2L	CHPF
MRE11A	KLHL24
PSRC1	PDGFRA
NUP88	DNAH10OS
RCC1	RP11-454H13.6.1
GEMIN4	ABCA7
C1orf74	TCP11L2
VRK1	S1PR1
PSMC1	HIST1H2BD
DOCK1	NEURL3
ALDH3A1	SYT5
GLT8D2	GPR35
KATNAL1	CARD9
SERBP1	CTD-2313N18.5.1
WDR77	LIPH
PHTF2	EFNA3
PSMB2	C3
BCAT1	MID2
STRA13	RP11-536G4.2.1
UBE2C	PNPLA7
PHEX	CYP7A1

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ADSL	TESK2
HSDL2	TNFSF15
CEP192	ZNF385C
GOT2	RP11-
	757G1.6.1
STIP1	FBXO2
VBP1	LRRN1
GNPNAT1	ZSCAN4
DLC1	PRSS27
FUBP1	EFNA1
HEATR1	CNNM1
TBC1D1	DDIT4
ECT2	SCN9A
GALNT1	C2
EXOG	CCDC114
CTD-	RRM2P3
2366F13.1.1	
POLA2	AP000696.2.1
SLC25A15	PDE4C
NEU3	CATSPERG
DOCK5	ELF5
CETN3	HS3ST3A1
BMP4	AC073321.5.1
CCDC72	CFB
SORL1	PARM1
UCHL5	C1orf145
ARPC4	C19orf51
FAM198B	CTC-
	454M9.1.1
KIAA1586	CCPG1
GSTO1	CTH
AC097359.1	MALAT1
BMP2K	TRIB3
SSBP1	NYAP1
C3orf67	KLF9
SMC3	LINC00176
AP1B1	IER3
ACAD9	GBP2
C2CD4C	VEGFA
SLIRP	SAT1
SNAPC1	BALAP3
RPL26L1	RP11-
	93B14.5.1
MGAT5B	ANKRD24
FSTL1	GAA
PARVB	CASP5
TIMM23	OASL
KLF12	MLXIPL
C1QBP	RP11-
	115C10.1.1
CCRL1	TSC22D3
SLC16A7	OPRL1
CDC6	SERPINA5
AC004381.6.1	SLCO4A1
CBFB	IGSF10
APAF1	S100P
PRKAR1B	EPAS1
FAM72D	SCARF1
FOXRED2	TNFSF10
TMEM209	ECT2L
CEP76	PSD
DLG5	CHAC1
AC003665.1.1	ANG
ACTL6A	CA9
PDP1	PIK3IP1
C11orf51	FLRT3
RPL39L	RP11-
	736K20.5.1
NUBPL	CEACAM22P
SFPQ	MYO15B
EME1	DPEP1
FANCM	KLK10
HMBS	ADM2
HNRNPR	ACCN3
HDAC9	LDB3
HECW1	FER1L4

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GCSH	CEBPD
LAPTM4B	TMEM105
GDF11	CCDC40
NAV2	RP3-
	395C13.1.1
NR2F2	ADCY4
INO80C	NKPD1
PPAT	SLC6A9
UMPS	C13orf33
CACNB4	SERPINA3
PLCL2	ACSS2
PRMT5	SLC2A6
ACADSB	EGR1
INCENP	TSPEAR-AS1
LARS2	MAPK15
ZNF714	RP11-
	178D12.1.1
CCDC86	FOS
PSMD2	ANGPTL1
LRRC20	ZNF467
ROCK2	RP11-
	712B9.2.1
C12orf48	PRSS35
TNPO1	NHLRC4
PPAP2B	SLC6A12
C14orf126	ICAM1
CDC42BPA	C2CD4A
BCCIP	GPR132
PSIP1	STRC
LSM3	C6orf223
MALT1	NFE2
AAK1	CTD-
	2319I12.1.1
CABYR	FAM167B
RBM12	PNCK
CYCS	TPPP3
SFMBT1	ITGA10
MPHOSPH9	ODF3B
SEH1L	WFDC10B
DCAF13	GRIP2
TIMM10	TXNIP
LYRM1	NUPR1
FAM171A1	ARRDC4
CENPH	SSPO
RP11-	TBX4
117L6.1.1	
BRIX1	LAMP3
PEG10	SLCO4C1
THOC7	HPN
CDCA7L	FAM132A
HPRT1	GAL3ST2
CEP170	NGFR
SF3A3	C2CD4B
PMVK	RANBP3L
HSPD1	C17orf28
TDP1	C21orf90
DNMT3B	MYO15A
CKAP2	LRRC4C
C12orf24	PLCH2
ANKRD28	HSPA6
CKAP5	IGFALS
POLR1E	KLB
DCUN1D5	AC005013.5.1
ORC1	CSF3R
KCTD1	RYR1
PSMA7	
CCDC134	
PSMC2	
PLS3	
PIGN	
CTD-	
2224J9.2.1	
NUP153	
ME3	
PITPNM3	
ABCD3	

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IQCC
DCLRE1B
USP1
AC108463.2.1
UPF2
SSRP1
ALG8
STARD13
NEK2
FCF1
GNG12
MASTL
TBC1D5
AP3B1
CBX3
PGRMC1
ATG3
POLH
PDCD11
RP11-
666A20.1.1
ZDHHC23
RGS17
FAM72A
SEPHS1
FAM122B
VOPP1
PSMA3
FLVCR2.1
LIN9
MANEA
FAM208B
GTPBP4
TTI1
CCDC88A
FAM48A
TMSB15B
SELRC1
RIMKLB
WDR17
ODZ3
CNOT1
SMCHD1
RDX
POLD3
LAS1L
CLTC
PPP1R14B
GEMIN5
AGK
C10orf125
SMS
PDAP1
LYPD6
RP11-
290F20.1.1
SLC36A1
TRAIP
RP6-
65G23.3.1
PPIA
RP11-
85G18.4.1
KCNC4
GDAP1
CABLES1
LGALS1
RPP30
PI4K2B
DSN1
ZNF620
USP6NL
C9orf100
CAV2
PTCD2
KIAA1147

-continued

TOP2B
TXNRD1
MYL6
RNASEH2A
RBFOX2
CCT6A
AMMECR1
DLD
BAI2
ORC5
TIMM8A
BEND6
TSC22D2
NAP1L5
ATIC
SIGLEC15
AP4S1
BAX
TTK
WDR12
TFAM
GPRIN1
PRDX1
GEMIN6
MMACHC
JUN
ERC1
EPHB2
XPO6
KIF24
SRGAP1
AURKB
TRRAP
GPD2
SCFD2
SMAP2
GSS

[0255] Supplementary Data 41: Comparison of overlaps between recurrent DE genes across all samples (vs. A38Per) and DMSO vs. 6AN treated A38Lg cells

Recurrent Up/ 6AN Down	Genes	Recurrent Down/ 6AN Up	Genes	Recurrent Up/ 6AN Up	Genes	Recurrent Down/ 6AN Down	Genes
Coregulated/ 6AN down: 255/1968 (13%)	PPIA	Coregulated/ 6AN up: 332/2192 (15%)	SAT1	Coregulated/ 6AN up: 82/2192 (0.04%)	EEF1A1	Coregulated/ 6AN down: 111/1968 (0.06%)	KRT19 + M2:M52
Coregulated/ Recurrent up: 255/891 (28.6%)	DYNC1H1	Coregulated/ Recurrent down: 332/1842 (18%)	ATP1B1	Coregulated/ Recurrent up: 82/891 (9%)	AKR1C2	Coregulated/ Recurrent down: 111/1842 (0.06%)	DCBLD2
	CCT5		GRN		RPL12		PLAU
	YBX1		JUP		RPL31		EZR
	HNRNPA3		LCN2		RPL29		GPR110
	XRCC6		CST1		IDI1		COL17A1
	PRDX1		TXNIP		XBP1		ANO1
	SERBP1		LAMB3		ATP2B1		PADI1
	DPYSL3		DDR1		HERPUD1		SDC1
	ECT2		DUOX2		MTHFD2		CD82
	CBX3		SLCO4A1		MAN1A1		CGN
	SLC25A3		GSN		SHMT2		LY6D
	HMGN2		LIMA1		MAP2		CRABP2
	TMPO		LYZ		ABCA3		DOCK9
	CYCS		MUC1		TOMM7		SPNS2
	PAICS		GNE		ASNS		RAPGEF3
	MYL6		DAB2IP		PYCR1		FGFBP1
	NOP56		TNIP1		TBC1D4		RBP1
	HNRNPR		LIPH		UCP2		PRSS12
	FKBP1A		RHBDF2		CELSR3		C1orf106
	THBS1		HS3ST1		LETMD1		QPCT
	DEK		RDH10		KLHDC2		STEAP2
	ASPM		MUC20		ARG2		ARHGDIB

-continued

NUP210	CLIP2	LEPRE1	ZNF185
PCNA	KLK10	SLC25A29	DHRS9
HNRNPAB	SLC44A2	PCK2	CDC42BPG
ATIC	EFNB1	SLC26A6	NBPF10
NCAPD2	NFKBIA	PTPRM	MUC4
RANGAP1	EVPL	SYT11	PLEKHA7
MYO1B	ENDOD1	FAM113A	AIG1
MTHFD1	ICAM1	GAB2	RP11-800A3.4.1
FAM208B	CDC42EP2	CRELD2	MMD
GPD2	CTSL1	SLC1A4	SAA1
GTPBP4	DRAM1	ABHD14B	ARAP2
TUBA4A	TRIM2	PDK1	ANKRD22
SMCHD1	NFKBIZ	C9orf150	CDS1
FOXM1	ADAM8	MLLT3	ALS2CL
SMARCC1	TRIB2	SSBP2	GALNT6
TFDP1	GSDMB	C3orf23	KRT15
XPO6	SGK1	DDT	XDH
TOP2B	ARHGEF16	SARM1	DGAT2
HEATR1	LRCH4	OSBPL6	FBP1
ODC1	HERC5	LEPREL2	CLDN10
ZWINT	PLEKHM1	EEF1A1P5	RHOD
ATXN10	ASAH1	HCN2	PRSS3
NUP155	TMEM8A	VLDLR	CTSL2
CDCA7L	PRICKLE4	BAMBI	SNAI2
SNRPB	NEU1	USE1	GALNT12
GNPNAT1	TNFSF15	CGREF1	P2RY2
WDR3	PLAUR	ATHL1	RAG1
HJURP	AC103810.2	GPR37	PADI3
KNTC1	RHBDF1	DERL3	TESC
REV3L	HSD17B11	PLK3	SELL
CDH2	ROBO4	SLC29A4	GPR116
FARSB	EREG	TMEM234	GCHFR
CCDC88A	VWA1	RP11-66N24.3.1	RP11-314P12.3.1
NETO2	TJP3	CORO6	CHST4
NUP153	PDCD4	C9orf37	SHANK2
USP1	PTPRE	EEF1A1P6	DNAH10
MCM6	C15orf48	DNAJC3-AS1	KRT23
POLD2	TPBG	C3orf78	COL1A1
SF3A3	TNFAIP3	MCEE	NMNAT2
TYMS	PROS1	RP11-480A16.1.1	AKAP6
KIAA1324	GPR108	ZNF70	AADAC
TMEM48	AGFG2	NAT6	VEPH1
TCOF1	RHBDD2	SEMA3F	C7orf58
PHIP	DGAT1	VKORC1	DNAH12
BRIX1	HBP1	RP5-1103G7.4.1	SLC22A20
UTP20	RNF103	P2RX6	MCC
CTPS	TSC22D3	C10orf102	RP11-7K24.3.1
RANBP1	CAPS	THTPA	ANKRD56
SACS	SLC2A6	FTX	CTD-2021J15.2.1
DTL	ABCA7	AC002472.8.1	ARL14
CDC20	TMC4	RP11-243J18.3.1	C15orf62
AP1B1	FAM193B	CTD-307407.5.1	METTTL7B
CDC6	VPS28	RP11-574K11.20.1	SLC4A3
ABCD3	ARSD	CES4A	KLHL13
EBNA1BP2	NCF2	METTTL12	RP11-357H14.19.1
PHTF2	MMP28	RP11-390P2.4.1	FGFR3
TRIP13	PARP10	RP11-216F19.1.1	PKDCC
RUVBL1	LZTS2	RP11-85K15.2.1	RP11-157P1.4.1
RCC1	ANKRD13D	SEPT7L	SMO
MYBL2	DNASE2		CD22
CHML	LYNX1		SYTL5
CCDC165	APOBEC3G		TNNT2
SEH1L	CDH3		ADAMTS14

-continued

WDR12	AQP3	ANXA10
TMEM56	NYNRIN	TTC3P1
ERC1	ABCG1	NMU
RFC3	SH3YL1	RP11-
		597D13.9.1
DEPDC1	FBXO2	UNC5C
RP1-	PPP1R16A	AADAT
239B22.1.1		
BMP2K	SESN3	GRHL1
NT5DC2	TMPRSS3	TMEM169
MPHOSPH9	HEATR7A	CYP24A1
HDAC9	KYNU	RP11-
		448G15.3.1
STMN3	METTL7A	ASB9
MRT04	FA2H	NAALADL2
CDCA8	ZC3H12A	FAM46B
S100A2	CFI	CFTR
PI4K2B	SLC15A3	C16orf74
MCM8	SIGIRR	RP11-
		416I2.1.1
GTF2H3	FER1L4	PLAC1
JUN	SLC39A11	ZNF658
TTF2	FOXQ1	SP6
FAM171A1	RASSF5	TTN
FKBP5	DHRS3	C8orf46
ADSL	ARHGEF3	GPR65
DIAPH3	S100P	LEF1
KIAA0586	ZG16B	RP11-
		6F2.4.1
POLQ	PTK6	DENND2A
EXO1	CDA	
EEF1E1	IL32	
ODZ3	ZNF862	
SSX2IP	CD14	
CEP97	TMED1	
ABCC4	G0S2	
RAD51AP1	ENDOV	
MRPL47	SMAD6	
C9orf140	MIA	
GEMIN5	CACNB1	
FANCD2	ARL4D	
CCNE2	JAK2	
UMPS	KCNMB4	
BARD1	IRAK2	
USP13	RELB	
CENPV	CEBPB	
SMTN	MAFF	
PRPS1	C9orf16	
CCNF	RALGPS1	
RFC5	LTB4R	
ATP5G1	CSF2RA	
POLR3G	C10orf32	
CBX2	SAT2	
NOP16	ALPPL2	
MCM10	CCDC69	
SLC16A7	NR1H3	
POP1	GRAMD1C	
CLSPN	FAM113B	
PFAS	WASH7P	
KLF12	TGFBR3	
GEMIN4	YPEL3	
C3orf26	LRRN1	
GMNN	MAST3	
CENPN	ABC7-	
	42389800N19.1.1	
KIF15	ZNF467	
GDAP1	XAF1	
THOC7	C16orf7	
OPN3	RP3-	
	395M20.8.1	
C22orf29	DNAH2	
BAG2	ADCY4	
ERCC2	CDC42EP5	
JAM3	ITGAX	
DERA	ANKRD42	
GPRIN1	NLRP1	

-continued

E2F2	AC021593.1
CDC45	N4BP2L1
ARRB2	CLDN15
NCS1	PIWIL4
SFMBT1	S1PR1
RAD54L	HLA-H
C1orf112	NEURL3
PRKAR1B	P2RX4
FANCM	AC007283.5.1
CENPL	CA11
SELRC1	DPEP1
DOCK10	TBC1D3F
ORC1	IER5L
ALDH1B1	PER1
BCL2	ZSWIM4
PSMC3IP	ITGA10
SNRNP25	IL15
PRIM1	SERPINA3
HSPA4L	RTP4
SUV39H2	GLI4
MAGOHB	C17orf103
HMGB1P5	SIX5
RP11-	SNPH
14N7.2.1	
WDR77	PIP5K1B
CENPH	TFEB
NME1	IDUA
LRRC8C	TMEM102
AC027612.6.1	SPRY3
METTL8	FMO5
POLR1E	MTRNR2L9
HOMER1	VSIG10L
RGS17	GIMAP2
DCLRE1B	PLCH2
POLE2	LRRC6
CENPW	LRFN3
SKA1	RILPL2
ADAT2	TNFSF12
CCDC41	CROCC
WDR4	HLA-K
CCDC18	PRICKLE3
PDSS1	CXCL3
C12orf24	NOS3
PARVB	FBXO6
MMP24	CCDC146
C14orf126	CSF2
SLC25A15	CEP85L
NEGR1	GEMIN8
CAP2	ZNF628
OIP5	CXCL2
KIF5C	ZDHHC1
ARHGAP11B	NR3C2
FAM86A	RP11-
	285F7.2.1
TBC1D7	PLEKHF1
CEP128	RP11-
	403I13.8.1
DLEU2	B3GALT4
BOLA3	NEIL1
COQ3	KNDC1
C17orf89	PON3
ARHGEF26	CFB
GDF11	ENDOU
ACTR3B	FOXO4
ASRGL1	DDO
XYLB	BBC3
RP1-	KB-
140K8.5.1	1460A1.5.1
SFXN2	SLC16A13
TIMM8A	GAS7
MNS1	RNF152
GTF2H2	ABCA6
AC009948.5.1	SLC16A6
ZNF239	RENBP
RP11-	KIAA1683
253E3.3.1	

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ASTN2	BBS12
CHAC2	CACNA2D4
CSMD3	UPK3B
RASSF2	AP001372.2.1
CCDC138	ZNF517
C1orf74	ACYP2
CCDC134	ARHGAP25
CABYR	PRX
RP6-	CES3
65G23.3.1	
PPIAP29	ACCN3
CYB5RL	C19orf51
RP11-	IL4I1
521B24.3.1	
RP1-	VMAC
152L7.5.1	
C20orf94	TNFRSF9
AL357673.1	RP11-
	757G1.6.1
ISPD	NFAM1
ELOVL2	RP11-
	353B9.1.1
RBM24	PDZD7
RP11-	CARD14
204J18.3.1	
DPF1	NYAP1
RP11-	CD34
381E24.1.1	
RP11-	RP11-
1334A24.4.1	325F22.3.1
MIR621	EXOC3L4
AC092329.1	RP5-
	1182A14.3.1
RP3-	JAK3
324O17.4.1	
CTD-	WDFY3-AS2
2574D22.2.1	
PDE1A	PDGFRA
RP11-	MLXIPL
33N16.1.1	
RP11-	KLHDC1
618G20.2.1	
	MIR29C
	RP11-
	712B9.2.1
	CTD-
	2292P10.4.1
	PCSK4
	CLDN9
	PODN
	COL11A2
	RP11-
	65J3.1.1
	PPP1R32
	EFHC2
	TMEM105
	AC005152.2.1
	CASP5
	TBX6
	DNAH10OS
	SCARF1
	RP11-
	420G6.4.1
	ROM1
	PYGM
	SLCO4C1
	CCDC114
	FAM71E1
	RP11-
	263K19.6.1
	CTD-
	2341M24.1.1
	PCDHAC1
	CLIP3
	C7orf63
	C17orf108

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ECEL1P2
LRRC24
ZNF385C
TPPP3
RP11-
108M9.4.1
ANKRD24
SRCRB4D
MIR29B2
DNAH7
RP11-
454H13.6.1
RP11-
369J21.5.1
TMPRSS5
PTPRCAP
BCO2
C2orf81
FAM66C
LINC00176
GPR132
SLC5A12
PDE4C
ICAM5
C20orf195
FBXO24
NGFR
TNFSF13B
MST1P9
CH25H
CTC-
523E23.1.1
NFE2
CTC-
378H22.2.1
FAM132A
GBP5
HOXD4
UCN
NKPD1
ELF5
LINC00324
RP11-
115C10.1.1
ANGPTL1
RP11-
536G4.2.1
FAM66D
RP4-
541C22.5.1
HSPA6
CTD-
2313N18.5.1
CTD-
2547L24.3.1

-continued

CYP7A1
IGFALS
RRM2P3
U7

[0256] Although the invention has been described with reference to the above example, it will be understood that modifications and variations are encompassed within the spirit and scope of the invention. Accordingly, the invention is limited only by the following claims.

SEQUENCE LISTING

Sequence total quantity: 33

SEQ ID NO: 1	moltype = DNA	length = 22	
FEATURE	Location/Qualifiers		
misc_feature	1..22		
	note = Synthetic		
misc_feature	1..22		
	note = Primer: Chr18_24.258F		
source	1..22		
	mol_type = other DNA		
	organism = synthetic construct		
SEQUENCE: 1			
gctcagccct gtatcagcca gc			22

SEQ ID NO: 2	moltype = DNA	length = 23	
FEATURE	Location/Qualifiers		
misc_feature	1..23		
	note = Synthetic		
misc_feature	1..23		
	note = Primer: Chr18_24.258R		
source	1..23		
	mol_type = other DNA		
	organism = synthetic construct		
SEQUENCE: 2			
gggttacagg tatgagccac tgc			23

SEQ ID NO: 3	moltype = DNA	length = 25	
FEATURE	Location/Qualifiers		
misc_feature	1..25		
	note = Synthetic		
misc_feature	1..25		
	note = Primer: Chr18_24.506F		
source	1..25		
	mol_type = other DNA		
	organism = synthetic construct		
SEQUENCE: 3			
aatggagaag tcaggaatgt agtcc			25

SEQ ID NO: 4	moltype = DNA	length = 27	
FEATURE	Location/Qualifiers		
misc_feature	1..27		
	note = Synthetic		
misc_feature	1..27		
	note = Primer: Chr18_24.506R		
source	1..27		
	mol_type = other DNA		
	organism = synthetic construct		
SEQUENCE: 4			
gtatttcatt tatcaagttg cagctcc			27

SEQ ID NO: 5	moltype = DNA	length = 25	
FEATURE	Location/Qualifiers		
misc_feature	1..25		
	note = Synthetic		
misc_feature	1..25		
	note = Primer: Chr18_24.834F		
source	1..25		
	mol_type = other DNA		
	organism = synthetic construct		
SEQUENCE: 5			

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tttgcttctc actccaagtt catcc		25
SEQ ID NO: 6	moltype = DNA length = 24	
FEATURE	Location/Qualifiers	
misc_feature	1..24	
	note = Synthetic	
misc_feature	1..24	
	note = Primer: Chr18_24.834R	
source	1..24	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 6		
caacctcagg aacaatgcat cagc		24
SEQ ID NO: 7	moltype = DNA length = 24	
FEATURE	Location/Qualifiers	
misc_feature	1..24	
	note = Synthetic	
misc_feature	1..24	
	note = Primer: Chr18_24.834R	
source	1..24	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 7		
caacctcagg aacaatgcat cagc		24
SEQ ID NO: 8	moltype = DNA length = 22	
FEATURE	Location/Qualifiers	
misc_feature	1..22	
	note = Synthetic	
misc_feature	1..22	
	note = Primer: Chr18_25.125.6F	
source	1..22	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 8		
cgaaacagtc cagctgctat gg		22
SEQ ID NO: 9	moltype = DNA length = 24	
FEATURE	Location/Qualifiers	
misc_feature	1..24	
	note = Synthetic	
misc_feature	1..24	
	note = Primer: Chr18_25.125.6R	
source	1..24	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 9		
cttggtatt gtgactggta ctgc		24
SEQ ID NO: 10	moltype = DNA length = 25	
FEATURE	Location/Qualifiers	
misc_feature	1..25	
	note = Synthetic	
misc_feature	1..25	
	note = Primer: Chr18_25.428.000F	
source	1..25	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 10		
ccaatgcact aatttaatgt catgc		25
SEQ ID NO: 11	moltype = DNA length = 25	
FEATURE	Location/Qualifiers	
misc_feature	1..25	
	note = Synthetic	
misc_feature	1..25	
	note = Primer: Chr18_25.428.000R	
source	1..25	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 11		
cgtgctaatt tctatggtag actgg		25
SEQ ID NO: 12	moltype = DNA length = 24	
FEATURE	Location/Qualifiers	

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misc_feature      1..24
                  note = Synthetic
misc_feature      1..24
                  note = Primer: Chr18_25.632F
source            1..24
                  mol_type = other DNA
                  organism = synthetic construct

SEQUENCE: 12
cctaataccaa tatgcctggt gtcc                                24

SEQ ID NO: 13      moltype = DNA length = 23
FEATURE            Location/Qualifiers
misc_feature      1..23
                  note = Synthetic
misc_feature      1..23
                  note = Primer: Chr18_25.632R
source            1..23
                  mol_type = other DNA
                  organism = synthetic construct

SEQUENCE: 13
ctggaagtct gagatcaagg tgc                                  23

SEQ ID NO: 14      moltype = DNA length = 27
FEATURE            Location/Qualifiers
misc_feature      1..27
                  note = Synthetic
misc_feature      1..27
                  note = Primer: Chr18_25.778F
source            1..27
                  mol_type = other DNA
                  organism = synthetic construct

SEQUENCE: 14
aataatcacg aagcacttct gtattgc                              27

SEQ ID NO: 15      moltype = DNA length = 26
FEATURE            Location/Qualifiers
misc_feature      1..26
                  note = Synthetic
misc_feature      1..26
                  note = Primer: Chr18_25.778R
source            1..26
                  mol_type = other DNA
                  organism = synthetic construct

SEQUENCE: 15
tcaccagcag acatagtcac acttcc                                26

SEQ ID NO: 16      moltype = DNA length = 24
FEATURE            Location/Qualifiers
misc_feature      1..24
                  note = Synthetic
misc_feature      1..24
                  note = Primer: Chr18_25.808F
source            1..24
                  mol_type = other DNA
                  organism = synthetic construct

SEQUENCE: 16
ccttgagggt ggagtctaca gagg                                  24

SEQ ID NO: 17      moltype = DNA length = 25
FEATURE            Location/Qualifiers
misc_feature      1..25
                  note = Synthetic
misc_feature      1..25
                  note = Primer: Chr18_25.808R
source            1..25
                  mol_type = other DNA
                  organism = synthetic construct

SEQUENCE: 17
ctgctagcgt agccatctga gatcg                                25

SEQ ID NO: 18      moltype = DNA length = 24
FEATURE            Location/Qualifiers
misc_feature      1..24
                  note = Synthetic
misc_feature      1..24
                  note = Primer: Chr3_25.398F

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-continued

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source                1..24
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 18
gccctgtctt cccagaatca ttgc                                24

SEQ ID NO: 19          moltype = DNA  length = 26
FEATURE               Location/Qualifiers
misc_feature           1..26
                      note = Synthetic
misc_feature           1..26
                      note = Primer: Chr3_25.398R
source                1..26
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 19
catgaagcct atgaagatca ttatgg                                26

SEQ ID NO: 20          moltype = DNA  length = 26
FEATURE               Location/Qualifiers
misc_feature           1..26
                      note = Synthetic
misc_feature           1..26
                      note = Primer: Chr3_25.540F
source                1..26
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 20
ttagccagc aagtattcta gcatgg                                26

SEQ ID NO: 21          moltype = DNA  length = 27
FEATURE               Location/Qualifiers
misc_feature           1..27
                      note = Synthetic
misc_feature           1..27
                      note = Primer: Chr3_25.540R
source                1..27
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 21
gtcagtgtga ttcagtaaca atgatgg                                27

SEQ ID NO: 22          moltype = DNA  length = 24
FEATURE               Location/Qualifiers
misc_feature           1..24
                      note = Synthetic
misc_feature           1..24
                      note = Primer: Chr3_25.622F
source                1..24
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 22
cctgctcaag gctgacatgt cacc                                24

SEQ ID NO: 23          moltype = DNA  length = 24
FEATURE               Location/Qualifiers
misc_feature           1..24
                      note = Synthetic
misc_feature           1..24
                      note = Primer: Chr3_25.622R
source                1..24
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 23
gtcggactcg atggtcagca ctgg                                24

SEQ ID NO: 24          moltype = DNA  length = 25
FEATURE               Location/Qualifiers
misc_feature           1..25
                      note = Synthetic
misc_feature           1..25
                      note = Primer: Chr3_25.733F
source                1..25
                      mol_type = other DNA
                      organism = synthetic construct

SEQUENCE: 24

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aacccgaaac tttcaatgca cttgg		25
SEQ ID NO: 25	moltype = DNA length = 25	
FEATURE	Location/Qualifiers	
misc_feature	1..25	
	note = Synthetic	
misc_feature	1..25	
	note = Primer: Chr3_25.733R	
source	1..25	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 25		
cttcctctat agtgaagacc ctagg		25
SEQ ID NO: 26	moltype = DNA length = 25	
FEATURE	Location/Qualifiers	
misc_feature	1..25	
	note = Synthetic	
misc_feature	1..25	
	note = Primer: Chr3_25.812F	
misc_feature	1..25	
	note = Primer: Chr3_25.733R	
source	1..25	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 26		
tatggccatt cttgcagcag taagg		25
SEQ ID NO: 27	moltype = DNA length = 26	
FEATURE	Location/Qualifiers	
misc_feature	1..26	
	note = Synthetic	
misc_feature	1..26	
	note = Primer: Chr3_25.812R	
source	1..26	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 27		
aaagttggct aaggacatga ataggc		26
SEQ ID NO: 28	moltype = DNA length = 25	
FEATURE	Location/Qualifiers	
misc_feature	1..25	
	note = Synthetic	
misc_feature	1..25	
	note = Primer: Chr3_25.973F	
source	1..25	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 28		
ggagattccc tcaggtgcct atacc		25
SEQ ID NO: 29	moltype = DNA length = 24	
FEATURE	Location/Qualifiers	
misc_feature	1..24	
	note = Synthetic	
misc_feature	1..24	
	note = Primer: Chr3_25.973R	
source	1..24	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 29		
ctggtgttcc aggcaccact gagg		24
SEQ ID NO: 30	moltype = DNA length = 20	
FEATURE	Location/Qualifiers	
misc_feature	1..20	
	note = Synthetic	
misc_feature	1..20	
	note = Primer: CDH2F	
source	1..20	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 30		
ttattactcc tgggtgcgagt		20

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SEQ ID NO: 31	moltype = DNA length = 20	
FEATURE	Location/Qualifiers	
misc_feature	1..20	
	note = Synthetic	
misc_feature	1..20	
	note = Primer: CDH2R	
source	1..20	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 31		
gagctgatga caaatagcgg		20
SEQ ID NO: 32	moltype = DNA length = 22	
FEATURE	Location/Qualifiers	
misc_feature	1..22	
	note = Synthetic	
misc_feature	1..22	
	note = Primer: TOP2BF	
source	1..22	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 32		
gttacaggtg gtcgtaatgg tt		22
SEQ ID NO: 33	moltype = DNA length = 22	
FEATURE	Location/Qualifiers	
misc_feature	1..22	
	note = Synthetic	
misc_feature	1..22	
	note = Primer: TOP2BR	
source	1..22	
	mol_type = other DNA	
	organism = synthetic construct	
SEQUENCE: 33		
ttggcttcag aagtcttcat ca		22

1-12. (canceled)

13. A method for identifying a subject's responsiveness to a treatment regime comprising analysis of changes in global chromatin modifications in a DNA sample from the subject prior to and following the treatment regime to provide a prognosis or analysis of responsiveness to the treatment regime.

14. A The method of claim 13, wherein identifying changes in global chromatin modification comprises identifying changes in large-organized heterochromatin lysine (K)-9 modified domains (LOCKS), changes in large LOCKs, and/or changes in euchromatin domains (ECDs).

15. The method of claim 14, wherein analyzing is by combinatorial methods.

16. (canceled)

17. The method of claim 15, wherein combinatorial methods comprise Western blotting, immunohistochemistry, chromatin immunoprecipitation (ChIP), whole genome bisulfite sequencing, or a combination thereof.

18. The method of claim 13, wherein the sample is DNA from a cancer cell from a solid tumor.

19. The method of claim 1, wherein the solid tumor is pancreatic ductal adenocarcinoma (PDAC).

20. The method of claim 14, wherein identifying changes in LOCKs comprises identifying changes in H3K9Me2 and/or H4K9Me3.

21. The method of claim 20, wherein detecting a loss of H3K9Me2 in LOCKs in a cell from a primary tumor or from a local metastasis is indicative of a likelihood of evolution to distant metastasis.

22. The method of claim 14, wherein identifying changes in ECDs comprises identifying changes in H3K26Me3 and/or H3K27Ac.

23. The method of claim 22, wherein detecting local reprogramming of H3K26Me3 and/or H3K27Ac over differentially expressed genes withing ECDs in a cell from a primary tumor or from a local metastasis is indicative of a likelihood of evolution to distant metastasis.

24. The method of claim 14, wherein a LOCK has a length ranging from about 100 kb to 30 Mb.

25. The method of claim 14, wherein a LOCK has an average length of about 232 Kb-311 Kb.

26. The method of claim 14, wherein a large LOCK has a length ranging from about 500 Kb to 30 Mb.

27. The method of claim 14, wherein a large LOCK has an average length of about 73 0 Kb-1,340 Kb.

28. The method of claim 14, wherein an ECD has a length ranging from about 50 kb to 12 Mb.

29. The method of claim 14, wherein a LOCK has an average length of about 207 Kb-277 Kb.

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