

## **Enhancer methylation dynamics contribute to cancer plasticity and patient mortality**

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## ABSTRACT

During development enhancers play pivotal roles in regulating gene expression programs; however, their involvement in cancer progression has not been fully characterized. We performed an integrative analysis of DNA methylation, RNA-seq and small RNA-seq profiles from thousands of patients, including 25 diverse primary malignancies and 7 body sites of metastatic melanoma. We found that enhancers are consistently the most differentially methylated regions (DMR) as cancer progress from normal to primary tumors and then to metastases, compared to other genomic features. Remarkably, identification of enhancer DMRs (eDMRs) enabled classification of primary tumors according to physiological organ systems and in metastasis eDMRs are the most correlated with patient outcome. To further understand eDMR role in cancer progression we developed a model to predict genes and microRNAs that are regulated by enhancer and not promotor methylation, which shows high accuracy with chromatin architecture methods and was experimentally validated. Interestingly, among all metastatic melanoma eDMRs the most correlated with patient survival were eDMRs that 'switched' their methylation patterns back and forth between normal, primary and metastases and target cancer drivers, e.g.: *KIT*. We further demonstrated that eDMR target genes were modulated in melanoma bone metastasis microenvironment, suggesting that eDMRs respond to microenvironmental cues in metastatic niches. Our findings that aberrant methylation in cancer cells mostly affects enhancers, which contribute to tumor progression and cancer cell plasticity, will facilitate development of epigenetic anticancer approaches.

## INTRODUCTION

The landscape of DNA methylation undergoes global changes in many cancers (Jones 2012) which contribute to genomic instability (Pogribny and Beland 2009), facilitate genetic mutations (You and Jones 2012), and alter gene expression programs (Easwaran et al. 2014). Most studies of DNA methylation changes in cancers have focused on promoter regions, since hypermethylation of promoters is a key mechanism for gene silencing (Esteller 2007). Less attention has been given to aberrant DNA methylation in other region of the genome, such as enhancers (Aran and Hellman 2013; Marzese et al. 2013; Ziller et al. 2013; Brocks et al. 2014), and to its influence on gene expression in cancer.

During development enhancers play pivotal roles in regulating expression programs (Hnisz et al. 2013), are characterized by tissue-specific chromatin marks (Heintzman et al. 2009), and their activation corresponds with nucleosome and methylation loss (Zhou et al. 2011; Buecker and Wysocka 2012; Taberlay et al. 2014). Recent studies have used these features to predict enhancer-gene interactions (Aran and Hellman 2013; Aran et al. 2013; Andersson et al. 2014; Aran and Hellman 2014; He et al. 2014); however, these methods provide limited insight into the roles of enhancers in cancer, in particular, the transcriptional consequences of abnormal enhancer methylation.

Cancer plasticity refers to the ability of tumor cells to transition between states and evolve under selective pressure, facilitating metastagenesis (Friedl and Alexander 2011; Tam and Weinberg 2013). Tumor heterogeneity and the microenvironment clearly impact cancer cell plasticity (Friedl and Alexander 2011; Brabletz 2012; Meacham and Morrison 2013). However, unlike genetic mutations, DNA methylation is a reversible modification, (Franchini et al. 2012), thus, we hypothesized that methylation changes could promote cancer progression by affecting cancer cell plasticity. To test this hypothesis, we extended the analysis of (Aran and Hellman

2013; Aran et al. 2013; Aran and Hellman 2014) and analyzed differential DNA methylation patterns in twenty five cancer types and seven sites of metastatic melanoma. Our analysis suggests that enhancer methylation changes can be indicative of patient outcome, and may contribute to cancer progression through cancer cell plasticity.

## RESULTS

### **Enhancers exhibit highly dynamic methylation patterns upon malignant transformation**

In order to methodically examine DNA methylation changes involved in malignant transformation we analyzed 23 distinct cancer types (overall 25 datasets). We used the genome-wide scale Illumina HumanMethylation450 arrays (Dedeurwaerder et al. 2011) to identify differentially methylated regions (DMRs) by analyzing over 6200 DNA methylation profiles of patients' tumor and normal tissue (Supplemental Table S1). We identified 123,649 DMRs (minimum region-wise mean methylation difference  $>0.3$ , false discovery rate,  $q < 0.2$ ). Interestingly, we found that enhancers had more DMRs than any other genomic region; promoters and CpG islands exhibited the less variation (Fig. 1A). This result suggests that alterations of enhancer methylation have a significant role in cancer progression, in support of data reported previously (Aran and Hellman 2013; Ziller et al. 2013; Aran and Hellman 2014; Taberlay et al. 2014). To test whether the selected threshold of 0.3 for differential methylation represented an optimal measure, we calculated the fraction of DMRs in each genomic feature using higher and lower thresholds (from 0.2 to 0.5); enhancers were consistently the most variable regions (Supplemental Fig. S1A). As the frequency of CpG dinucleotides varies with genomic features (Supplemental Fig. S1B), we repeated the differential methylation analysis using methylation levels of individual CpG sites in the array (Supplemental Fig. S1C), and observe consistent results, namely that CpG

methylation within enhancers are the most variable compared to all other genomic features.

Given the parallels between embryonic development and oncogenic transformation (Hon et al. 2013), we examined the binding patterns of central pluripotent transcription factors (TFs) *POU5F1*, *SOX2*, and *NANOG* in various DMRs. We observed that multiple binding occurred more frequently at differentially methylated enhancers than at promoters or CpG islands (Fig. 1B). These results support previous findings (Göke et al. 2011; Whyte et al. 2013) suggesting that eDMRs bound by pluripotent TFs may mediate central expression programs. To qualitatively examine the altered enhancers, we analyzed information on ChromHMM states (Ernst and Kellis 2012). We found that compared to enhancers that were not differentially methylated (static enhancers), eDMRs were significantly enriched for the ChromHMM state of 'strong enhancers' ( $P < 2e-16$ , Fig. 1C), similar to previous reports (Aran et al. 2013). This results suggest that enhancers altered in cancer, may regulate their target genes more strongly than do unaltered enhancers. It is known that promoter and island hypermethylation are associated with gene silencing in cancers (Bergman and Cedar 2013), and, indeed, we found that these regions were frequently hypermethylated (63% and 94%, respectively). In comparison, enhancers were mostly differentially hypomethylated (67%, Fig. 1D). Next, used chromatin marks to qualitatively determine the chromatin features of the hypermethylated and hypomethylated enhancers. Using data from the ENCODE Project (Rosenbloom et al. 2013), we found that differentially hypomethylated enhancers exhibited significantly higher levels of chromatin marks of active DNA (increased DNase I, H3K4me1, H3K4me2, H3K27ac, H2A.Z, EP300, POL2, and decreased CTCF). Whereas, differentially hypermethylated enhancers showed the footprints of closed and inactive DNA (Fig. 1E and Supplemental Fig. S1D-E), consistent with previous results (Zhou et al. 2011; Aran et al. 2013; Lam et al. 2014). Taken together, the

analysis of multiple tumor types strongly suggests that alteration of enhancer methylation is a frequent feature of transformed cells, and that it generally leads to chromatin activation, likely involved in cancer progression.

### **Differential methylation patterns of eDMRs clusters tumors according to their organ system**

Next, we compared the patterns of methylation change in enhancers between the various types of cancer. Since enhancers show tissue-specific patterns of histone modifications and TF binding in normal cells (Bulger and Groudine 2011), we expected that alteration of methylation patterns would be cancer/tissue-specific. Indeed, most eDMRs (54%) were unique to a single cancer type (Fig. 2A). Strikingly, principal components analysis (PCA) of eDMRs showed a higher order clustering of cancers into groups related to the same organ system (Fig. 2B). For example, the central nervous system cluster includes cancers arising from glia and astrocytes (Fig. 2B, green circle); similarly, the reproductive system cluster consists of breast, uterine, and prostate tumors (Fig. 2B, pink circle). Importantly, PCA analyses of differentially methylated islands, promoters, and intergenic regions showed no such clustering (Supplemental Fig. S2A-C). No library preparation batch effect was found, either (Supplemental Fig. S2D-E). This evidence provides confidence that our findings were not affected by batch effect, and strengthen our hypothesis that enhancer methylation changes may have meaningful biological roles.

Next, in order to uncover the functional effect of altered enhancer methylation, we first compared the genomic neighborhood of eDMRs with static enhancers (those showing no differential methylation). We found that eDMRs were flanked by more genes and microRNA (miRNA) than were static enhancers (Supplemental Fig. S2F-G). To identify which genes the eDMRs regulate, we developed an integrated model

combining multi-omics data for associating enhancers with their target genes (eDMR-associated genes) (Fig. 2C and Supplemental Fig. S2H for pipeline). Since it is known that there is an inverse correlation between DNA methylation and chromatin activity (Zhou et al. 2011), we retained only inversely correlated eDMR-gene pairs, which represented the majority of our data (Supplemental Fig. S2I). In order to assess the validity of our eDMR-gene pair predictions, we compared our results to other methods that identify enhancer-promotor associations based on physical interactions IM-PET (He et al. 2014), ChIA-PET, Hi-C (Teng et al. 2015), and transcriptional activities of interacting enhancer-promoters (cap analysis gene expression. CAGE) (Andersson et al. 2014). Our model predicted eDMR-gene pairs separated by 400 kb or less at a precision rate of 75% or better (Fig. 2D), indicating that our model can reliably predict interacting enhancer gene pairs.

Similar to their enhancers (Fig. 2B), the genes predicted to be regulated by the eDMRs, also showed organ system patterns of enrichment (Fig. 2E, Supplemental Table S3). Using our model we were identify known tissue-specific and ubiquitous oncogenic genes and microRNAs (miRNAs) (Fig. 2F-H), such as: *ESR1* in breast cancer (Holst et al. 2012; Aran and Hellman 2014), *ECT2* in lung cancer (Murata et al. 2014), and *WNT3A* in prostate cancer (Verras et al. 2004), hsa-miR-9-1 (Ma et al. 2010) and the clustered miRNAs hsa-miR-200a and miR-200b (Korpál et al. 2008) (see Supplemental Table S4 for full list of predicted eDMR-gene pairs). Taken together, our model defines cancer-related eDMRs as organ-system-specific regulators of genes and miRNAs that are central to malignant transformation.

### **Accumulation of eDMR hypomethylation correlates with likelihood of metastasis**

Our analysis showed that changes in methylation patterns of enhancers in primary tumor tissues are influenced by their relation to the organ system (Fig. 2B), we were curious what would occur to DNA methylation patterns of cells that disseminate from the primary location, and colonize in metastatic sites. To evaluate this, we focused on melanoma, a highly metastatic cancer (Braeuer et al. 2014). We compared methylation data from metastatic melanoma patients (7 distinct locations) to data from patients with primary *in situ* melanoma (Supplemental Table S1). Consistent with our observation of differential methylation patterns in primary tumors (Fig. 1A), the majority of 'metastatic DMRs' occurred within enhancers (Fig. 3A), suggesting that enhancers play important roles, not only in malignant transformation, but also in metastatic progression. Additionally, metastatic eDMRs much better differentiate between patient outcomes than do DMRs from any other genomic feature (Fig. 3B and Supplemental Fig. S3A). The majority of eDMRs were specific to a single metastatic site (Fig. 3C and Supplemental Fig. S3B), and were mostly (74%) hypomethylated (Fig. 3D), similar to our observations in primary cancers (see Fig. 2A and Fig. 1D). Strikingly, Fig. 3E shows that accumulation of hypomethylated enhancers highly correlates with the likelihood of forming metastases at distant organs (derived from (Meyers and Balch 1998)). For example, melanoma is more likely to metastasize to lymph nodes than to the brain, correspondingly, a greater number of enhancers are differentially hypomethylated in brain metastases than in lymph node metastases (2.24% and 0.33%, respectively; Fig. 3E). In contrast, we found no correlation between the fraction of differentially hypermethylated metastatic eDMRs and the frequency of spreading to organs (Supplemental Fig. S3C). These results are in agreement with studies suggesting that global hypomethylation is a common feature of diseased states (Pogribny and Beland 2009). Since widespread DNA methylation changes are associated with aging (Richardson 2003), we calculated the correlations between patient ages and accumulation of enhancers methylation changes. Encouragingly, we found no significant correlation between

eDMR methylation changes in metastatic tissues and the patient age (Supplemental Fig. S3D), suggesting that metastatic progression corresponds to cancer-related changes, and not age-related changes.

In order to specifically explore whether melanoma eDMRs could promote metastatic growth, we identified genes differentially expressed between primary and metastatic melanoma, and compared the correlation of these genes with methylation patterns of differentially methylated enhancers and promoters (eDMRs and pDMRs, respectively). We found that eDMRs were significantly more correlated with differentially expressed genes than were pDMRs (Supplemental Fig. S3E). Next, we divided the genes into two groups: eDMR-associated genes and all other genes not identified by our model to be associated with eDMRs (control genes, Fig. 3F), and examined the differential expression patterns of these two groups. The percent of genes that were differentially expressed between normal and primary melanoma was similar for both groups (60% and 53% for eDMR-associated genes and control genes, respectively; Fig. 3F), however, between primary and metastatic melanoma the eDMR-associated genes were significantly more variable than the control genes (binomial distribution,  $P < 2e-16$ ; 57% and 3% respectively, Fig. 3F, marked by gray areas). This observation supports our notion that a functional interaction exists between the eDMRs and their predicted associated genes. Additionally, we note that a fraction of the control genes were downregulated in both primary and metastatic melanoma (Fig. 3F, left panel, marked with dashed box). This group was enriched ( $q < 0.01$ , FDR corrected) with Gene Ontology (GO) terms associated with tissue development (GO:0008544, GO:0007398) and differentiation (GO:0030216, GO:0009913, GO:0030855), likely a feature of tumor de-differentiation (Brabletz 2012).

## **Methylation plasticity of melanoma eDMRs is associated with increased patient mortality**

The transition of cancer from the *in situ* primary stage to the metastatic stage involves phenotypic plasticity (Craene and Berx 2013), which facilitate migration through tissues and adaptation to changing microenvironments. Underlying tumor plasticity are genetic and epigenetic regulatory layers that are reprogrammed in the context of cancer (Friedl and Alexander 2011). Interestingly, we found that 18% (N=277) of melanoma metastatic eDMRs (N=1539) switched the direction of methylation change as melanoma progressed (denoted 'switched' eDMRs; Fig. 4A, *bottom* circle). For example, regions that were hypermethylated between normal melanocytes and primary melanoma, were hypomethylated between primary and metastatic melanoma. We also defined two other eDMRs groups: 'consistent' and 'de novo'. The 'consistent' eDMRs exhibited differential methylation changes in the same direction (e.g., hypomethylated between normal melanocytes and primary melanoma, and hypomethylated between primary and metastatic melanoma; Fig. 4A, *right* circle). The 'de novo' group exhibited differential methylation only between primary and metastatic melanoma samples, and not between normal melanocytes and primary tumors (Fig. 4A, *left* circle). We examined whether these three groups of eDMR differed with respect to metastatic progression, by comparing their ability to correlate with survival of patients (see Methods). To our knowledge, this is one of the first examples (Stone et al. 2015) of a survival analysis applied to DNA methylation patterns of enhancers. We identified 30 eDMRs that were associated with patient survival rates (marked 'survival' eDMRs, Fig. 4A, *middle* circle). Remarkably, these survival eDMRs were enriched with the switched eDMRs (Fig. 4A), whereas de novo eDMRs were significantly depleted (Fig. 4A). In addition, the switched eDMRs exhibited the highest conservation scores (Fig. 4B), and the fewest copy number variations (CNVs) in melanoma patients (Fig. 4C). These results suggest that

eDMRs, in particular the switched eDMRs that exhibit methylation plasticity, are functionally important in cancer.

Given these results, we hypothesized that methylation plasticity may play important roles in melanoma metastatic progression. To examine this, we performed survival analyses also the eDMR-associated genes, using their expression patterns across patients. Remarkably, we found that 40% were able to significantly differentiate between survival outcomes (Chi-square,  $q < 0.1$ , FDR corrected); these genes include: *ATP2B1* (Lee et al. 2002), *FMNL2* (Zhu et al. 2008), *KIT* (Tian et al. 1999), *PRKCE* (Sharif and Sharif 1999), and *VGF* (Mitra et al. 2008). Figure 4D represents such an example in which the oncogene *KIT* and the eDMR (located at Chr4:55708295-55709294, hg19/GRCh37) not only exhibit switched patterns of methylation and expression, but both independently distinguish between patient survival rates based on their expression and methylation patterns, respectively (Fig. 4D, *left* and *right* panels). The *KIT* promoter was not differentially methylated, nor did the methylation pattern differentiate between survival times (Fig. 4D, *middle* panel).

To experimentally test the role of enhancer methylation on the transcriptional regulation of *KIT*, we first examined *KIT* expression upon treatment with 5-aza-2'-deoxycytidine (5-aza-dC), a DNA methyltransferase inhibitor. There was a significant decrease in *KIT* levels in treated compared with untreated cells (Fig. 4E). Next, we cloned the *KIT* enhancer upstream of luciferase reporter, and demonstrated that its expression was reduced upon enhancer methylation (Fig. 4F and Supplemental Fig. S4A). Finally, we examined the role of *KIT* upregulation on the invasive potential of melanoma cells (Fig. 4D). Remarkably, noninvasive melanoma cells (Golan et al. 2015) acquired significant invasion ability upon *KIT* over-expression (Fig. 4G and Supplemental Fig. 4B). Taken together, our data demonstrate that enhancer methylation contributes to cancer progression by directly regulating oncogene expression.

Next, we asked whether the dynamics of eDMR methylation and eDMR-associated genes expression are a result of tumor heterogeneity, evolution of the disseminated cancer cells, or are induced by the metastatic niche. To assess this, we selected two eDMR-associated genes, *CYTL1* and *KIF14*, which were differentially expressed only between primary and metastatic melanoma (not between normal and primary melanoma tissues). *CYTL1* is a cytokine-like protein implicated in lung cancer and neuroblastoma (Kwon et al. 2012; Wen et al. 2012), and *KIF14* is an oncogene essential for cytokinesis (Corson et al. 2005); both are upregulated in melanoma bone metastasis ( $P < 0.05$ , Fig. 5A, *top* and *bottom* panels). High expression levels of these genes is correlated with poor patient survival rates (Fig. 5A, green stepwise curves; Chi-square  $q < 0.1$ , FDR corrected). Enhancers of these genes were differentially methylated, yet their promoters were not (Fig. 5A *middle* and *right* panels, respectively). To experimentally analyze the role of enhancer methylation on the transcriptional regulation of *CYTL1* and *KIF14*, we first examined whether their expression is sensitive to DNA methylation. *CYTL1* and *KIF14* expression was significantly upregulated upon treatment of cells with 5-aza-dC (Fig. 5B). Next, we cloned the identified enhancers of *CYTL1* and *KIF14* upstream to a luciferase reporter, and observed a decrease in the reporter expression upon enhancer methylation (Fig. 5C and Supplemental Fig. S4A). These results strengthen the validity of our model by demonstrating that the identified eDMRs, elicit response to changes in their methylation and directly regulate the expression of their associated genes: *KIT*, *CYTL1*, and *KIF14*. Finally, to assess changes in gene expression induced by metastatic melanoma cells colonization in the bone tissue, we first generated a melanoma cell line stably expressing the *GFP* gene, to enable tracking of the melanoma cells. Next, we established a co-culture of melanoma cells with human primary osteoblasts (Dillon et al. 2012) (Fig. 5D), and found that levels of *CYTL1* and *KIF14* were significantly increased, compared to their levels in melanoma cells cultured alone (Fig. 5E). Our data support the notion that homing into a new

microenvironment induces expression of pro-cancer genes, which are regulated by eDMRs in melanoma metastatic cells. Taken together, our data suggest that methylation changes at enhancers contribute to melanoma phenotypic plasticity and ultimately to the patient's chance of survival.

## DISCUSSION

Here we performed an analysis of DNA methylation alterations in over 6200 cancer patients from thirty-one cancer datasets, including twenty-three diverse primary malignant tumors, two benign tumors, and melanoma metastases to seven distinct organs. This extensive analysis revealed that most methylation variation occurs at enhancers (Fig. 1A, Fig. 3A and Supplemental Fig. S1C). Changes in methylation patterns could result from competition between methylation and de-methylation processes or from errors in replication (Jones 2012). Our findings support the latter hypothesis, for two reasons. First, replication-related methylation errors occur more frequently in methylated regions, such as enhancers, where replication errors lead to loss of methylation; this is what we observe (Fig. 1D). Second, expression of *DNMT3B*, which encodes a methylase enzyme was upregulated in many tumors, and hence, we would expect increased methylation, since we observed no significant differences in expression of *TET* genes, which encode demethylase enzymes (Supplemental Fig. S2J). However, since the majority changes in enhancers' methylation involves demethylation, we hypothesize that loss of enhancer methylation could have occurred during replication.

Enhancers play central roles in normal development and differentiation by responding to complex environmental cues. Cancer cells are exposed to changing environmental conditions that requires their adaptation; this mainly occurs through epigenetic reprogramming (Friedl and Alexander 2011; Goding et al. 2014). We hypothesize that in the context of cancer, enhancer methylation may be primed to

respond to microenvironmental signals. To test this, we mimicked the microenvironment of melanoma bone metastases, and recapitulated the perturbation of expression of eDMR-associated genes that were altered in the bone metastases. These results suggest that the metastatic niche can alter expression of eDMR-associated genes.

Interestingly, we show that in comparison to DMRs in other genomic features (islands, promoter, exons, introns, etc.) eDMRs can differentiate best between patient outcomes (Supplemental Fig. S4D). These results demonstrate the important role of alteration of enhancer methylation in cancer progression. However we do not exclude that other factors may drive cancer progression, nor are we suggesting a direct causality between enhancer methylation and cancer progression, it may well be that alteration of enhancer methylation is a secondary event of the malignancy.

Nevertheless, we did observe that methylation patterns of eDMRs may be informative of patient survival rates (Fig. 3B). Within this group, was a subset (18%) of highly conserved eDMRs that displayed methylation plasticity ('switched' eDMRs, Fig. 4A-C), and provided insight into alteration of their associated genes (Fig. 4D). This is one of the first studies (Stone et al. 2015) to suggest that methylation patterns of enhancers can be used to predict patient outcome. Moreover, it has been shown that in many diseases, including cancer, methylation changes are accumulative as the cancer progresses (Pogribny and Beland 2009); here we show that methylation plasticity may also play important roles in cancer progression (see Supplemental Fig. S4C for model of methylation plasticity and cancer progression). A prime example of the relationships between metastatic progression, methylation plasticity, and patient mortality is that of the eDMR-associated oncogene, *KIT* (Fig. 4D-F). Both the methylation of the eDMR and the expression of the *KIT* are display plasticity (Fig. 4D), and significantly distinguish between patient survival rates. Our results suggest

that enhancer methylation patterns may be informative of patient outcomes, and that they may influence malignant progression via methylome plasticity.

## FIGURE LEGENDS

### Figure 1

#### **Enhancers exhibit highly dynamic methylation patterns upon malignant transformation**

(A) Heatmap shows that across 25 cancer datasets (x-axis) differential methylation occurs mostly within enhancer regions. Colors indicate high (pink) to low (green) relative amount of DMRs in Illumina HumanMethylation450 arrays (see Supplemental Fig. S1C for differentially methylated CpG sites, and Fig 3A for regions differentially methylated between primary and metastatic cancer). (B) ChIP-seq signals of pluripotency transcription factors (TFs): *POU5F1*, *SOX2*, and *NANOG* are higher within differentially methylated enhancers (eDMRs) than within differentially methylated islands or promoters; average peaks across all cancer datasets are shown. (C) 'Strong enhancer' ChromHMM chromatin annotation state (see Supplemental Table S2 for cell lines) is more enriched within differentially methylated enhancers ('eDMRs', orange area) than within enhancers that are not differentially methylated ('static enhancers', gray area). (D) Direction of methylation change between normal tissue (N) and primary cancer (P) in differentially methylated enhancers, islands, and promoters. Coloring indicates high (blue) to low methylation levels (yellow); heatmap heights are relative to amount of the differentially methylated enhancers. (E) Hypomethylated eDMRs exhibit footprints of open chromatin marks: increased DNase I hypersensitivity, increased POL2 and EP300 occupancy, increased H2A.Z, H3K4me1, H3K4me2 and H3K27ac marks, and decreased CTCF

binding. In contrast, hypermethylated eDMRs exhibit footprints of closed chromatin. *P*-values were calculated with two sample Student's *t*-tests and FDR adjusted. See Supplemental Fig. S1D-E for chromatin marks in other cell states and Supplemental Table S2 for cell lines information. All figures refer to DMRs determined between normal samples and primary tumors.

## Figure 2

### Variation in enhancer methylation classifies primary tumors according to their organ systems

(A) Pie chart shows that most eDMRs are unique (77% of the eDMRs appear in only one or two cancers; excluding the same cancer type, e.g., GBM1 and GBM2). (B) Principal components analysis (PCA) shows that differential methylation patterns of eDMRs are similar for cancers belonging to the same organ systems (circled and colored similarly). First and second principal components (x-axis: PC1; y-axis: PC2) account for 72% of variation (see Supplemental Fig. S2A-C for PCA analysis of methylation changes within islands, promoters and intergenic regions). (C) Schematic of method for identification of eDMR-gene and eDMR-miRNA pairs (see Supplemental Fig. S2H for flowchart). (D) *Top*: Precision of eDMR-gene pair predictions is compared to predictions from CAGE (Andersson et al. 2014), IM-PET (Teng et al. 2015), ChIA-PET, and Hi-C (Teng et al. 2015) methods (see text). *Bottom*: Percent of predicted eDMR-gene pairs. Results in both plots are divided into groups based on the distances between methylation site and transcription start site. Most predictions (>56%) lie within a distance of <500 kb (abbreviations: 100K, 0-100,000 bp, 200K: 100,001-200,000 bp, and so forth). (E) EDMR-associated genes are distinctly enriched for genes involved in diseases related to a particular organ system. Disease-related genes were derived from the DISEASES database (see

Methods). Scores represent  $-\log_{10}(\text{FDR corrected } P\text{-values})$ . (F) Scatter plots (a point for each patient) show examples of tissue-specific genes linked to cancer risk and their eDMRs (the same six cancer types shown in panel E). Spearman's correlations between eDMRs methylation and associated-gene expression are shown. (G) Landscape of two differentially hypomethylated eDMRs in uterine (UCEC) and head and neck (HNSC) cancers, associated with a single miRNA, miR-9-1, which is upregulated in both cancers. (H) Landscape of two differentially hypomethylated eDMRs in breast (BRCA) and colon (COAD) cancers associated with two miRNAs of the same family, miR-200a and miR-200b; each is upregulated in the respective tumors. In figures G -H, yellow arrows mark hypomethylated eDMRs, green arrows mark upregulated miRNAs; gray boxes mark eDMR and miRNA regions in the hg19/GRCh37 genome; distances between miRNAs and eDMRs are indicated on arrows. All figures refer to DMRs determined between normal samples and primary tumors.

### Figure 3

#### **Accumulation of eDMR hypomethylation correlates with likelihood of metastasis and patient outcome**

(A) Heatmap shows that enhancers are more differentially methylated than other genomic features (y-axis; sorted according to fraction of DMRs in each category; see Fig. 1A). (B) Enhancers are significantly more enriched with DMRs that can differentiate between patient survival times (see Methoda). Green colors represent enrichment and red depletion, respectively. Y-axis represents the  $-\log_{10}(\text{FDR corrected } P\text{-values})$  significance (binomial distribution). (C) Most melanoma eDMRs (70%) are exclusive to one metastatic site (see Figure 2A). (D) Heatmaps shows that in most eDMRs are hypomethylated between primary and metastatic melanoma

(74%). (E) Accumulation of eDMR hypomethylation is negatively correlated with the likelihood of forming melanoma metastases at each body location (Pearson's correlation,  $r=-0.71$ ,  $P=0.039$ , one-sided hypothesis testing). (F) Compared to all other genes (control genes; y-axis, *right* panel), genes associated with eDMRs (y-axis, *left* panel) are differentially expressed between primary melanoma and metastatic melanoma (57% and 3%, respectively; marked by gray areas). Whereas, differential expression between melanocytes and primary melanoma is similar for both groups (control genes, 53%, *right* panel, eDMR-associated genes 60%, *left* panel, x-axes). Abbreviations: adrenal, metastases to adrenal glands; gastro, metastases to the gastrointestinal tract; lymph, metastases to lymph nodes; and subcut, sub-cutaneous metastases. All figures refer to differential methylation between patients with primary melanoma and patients with metastatic melanoma.

#### Figure 4

##### Methylation plasticity of melanoma eDMRs is associated with patient mortality

(A) Venn diagram of three types of eDMRs found in metastatic melanoma: 'de novo', 'consistent', and 'switched' (described in text). Heatmaps display differential methylation between normal melanocytes and primary melanoma (N->P), and between primary and metastatic melanoma (P->M). 'Switched' eDMRs are enriched for 'survival' eDMRs and 'de novo' eDMRs are depleted (hypergeometric distribution, \*,  $P=9.4e-5$ ; \*\*,  $P=1.6e-2$ , respectively). Numbers represent amount of eDMRs in each category. (B) PhastCons conservation scores of enhancer average conservation (See Methods). 'Switched' eDMRs have the highest conservation (two-way ANOVA analysis, \*,  $P=4.2e-6$ , \*\*,  $P=9.9e-10$ , \*\*\* $P=2.9e-11$ ); bars represent average sequence conservation, error bars represent standard deviation. (C) Copy number variations (CNVs) are infrequent within eDMRs compared to static enhancers (two-way ANOVA analysis, \*\*\*,  $P<2e-16$ ). (D) *Top panel*: Schematic representation of the eDMR-gene pair: Chr4:55708295-55709294 and *KIT* gene. The

eDMR (pink) is interacting with the promoter (cyan) of *KIT* gene (green) through chromatin looping. The distance between the eDMR and TSS. *Bottom panel: Left:* Kaplan-Meier survival plot shows significant differences between survival times of patients based on the methylation levels of the eDMR (located at Chr4:55708295-55709294) of the gene *KIT* (Chi-squared, \*\*\*\*,  $P=2.9e-4$ ). Patients were divided into two groups of higher (blue) and lower (yellow) levels compared to the median (see Methods). *Middle:* Plot shows that patients with higher and lower methylation of the *KIT* promoter show no significant difference in survival rates. *Right:* Plot shows significant differences between survival times of patients with higher (green) and lower (red) expression levels of *KIT* (Chi-squared, \*\*\*,  $P=2.4e-3$ ). *Insets: Left and Right insets:* Both eDMR methylation and *KIT* expression exhibit switched patterns between normal melanocytes (N), primary melanoma (P) and metastatic melanoma (M). *Middle inset:* *KIT* promoter does not exhibit switched methylation patterns.  $q<0.05$ , \*,  $q<0.01$ , \*\*. (E) Inhibition of DNA methylation increases *KIT* expression. WM3682 melanoma cells were treated with 10  $\mu\text{m}$  5-aza-dC or with DMSO (control), *KIT* mRNA expression was normalized to levels of *GAPDH*. Data are relative to levels in control treated cells. Error bars represent  $\pm$  SEM,  $P<0.05$ , \*; (N=3). (F) WM3682 melanoma cells were transfected with methylated *KIT*-eDMR reporter plasmid, unmethylated *KIT*-eDMR reporter, or reporter without eDMR (control). Firefly luciferase activity was normalized to *Renilla* luciferase activity. Fold-changes are relative to control. Error bars represent  $\pm$  SEM;  $P<0.05$ , \*; (N=3). (G) *KIT* overexpression increases melanoma invasiveness. WM3682 melanoma cells were transfected with *KIT* expression vector or empty vector (control), invasion ability was analyzed. Error bars represent  $\pm$  SEM,  $P<0.05$ , \*; (N=3). *Bottom panel:* a representative image of invading cells.

**Figure 5**

## Melanoma co-culture with osteoblasts alters pro-cancer eDMR-associated gene expression

(A) Schematic representation of the eDMR-gene pairs: Chr4:5019792-5020791 for *CYTL1* and Chr1:201198504-201199503 for *KIF14*. The eDMR (pink) is interacting with the promoter (cyan) of *KIT* gene (green) through chromatin looping. Kaplan–Meier survival plots (*left panels*) show significant differences between patient outcomes at high and low expression levels (green and red curves, respectively) of *CYTL1* and *KIF14* genes. *Insets*: Bar graphs show upregulation of *CYTL1* and *KIF14* in melanoma bone metastases. *Middle panel*: Methylation of the promoters of these genes does not vary between primary melanoma (P) and melanoma bone metastases (M). *Right panel*: Enhancers are differentially hypomethylated in melanoma bone metastases (Wilcoxon rank-sum tests, \*,  $q < 0.15$ , \*\*,  $q < 0.05$ ; FDR adjusted). (B) WM3682 melanoma cells were treated with 10  $\mu\text{m}$  5-aza-dC or with DMSO (control) followed by quantification of *CYTL1* and *KIF14* mRNA levels that were normalized to *GAPDH*. Data are relative to levels in control-treated cells. Error bars represent  $\pm$  SEM, \*,  $P < 0.05$  (N=3). (C) WM3682 melanoma cells were transfected with methylated *CYTL1*-eDMR or *KIF14*-eDMR reporter plasmids, unmethylated eDMR reporter or reporter without eDMR (control). Firefly luciferase activity is normalized to *Renilla* luciferase. Fold-changes are relative to control, error bars represent  $\pm$  SEM, \*,  $P < 0.05$  (N=3). (D) Experimental design scheme. (E) Levels of *CYTL1* and *KIF14* mRNA were determined in melanoma cells before and after co-culturing with osteoblasts. Data were normalized to levels of *actin*. Error bars represent  $\pm$  SEM, \*,  $P < 0.05$  (N=3).

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## **METHODS**

### ***TCGA and GEO patients data sources: DNA methylation, RNA-seq, smRNA-seq***

Publically available data of DNA methylation, RNA-seq gene expression, and small RNA-seq miRNA expression from cancer patient tissues were obtained from The Cancer Genome Atlas (TCGA, <https://tcga-data.nci.nih.gov/tcga>) and from the National Center for Biotechnology Information (NCBI) Gene Expression Omnibus (GEO) repository (<http://www.ncbi.nlm.nih.gov/gds>). See Supplemental Table S1 dataset information.

### ***Chromatin marks, transcription factors (TFs), sequence conservation and copy number variation data***

For analysis of chromatin marks (Fig. 1E and Supplemental Fig. 1D-E), we downloaded RNA polymerase II (POL2), histone modifications: H2A.Z, H3K4me1, H3K4me2 and H3K27ac, histone acetyltransferase (EP300), CCCTC-binding factor (CTCF), DNase I hypersensitivity and Chromatin State Segmentation by HMM (ChromHMM) from the ENCODE Project. Replicate experiments were averaged. See Supplemental Materials for URL and description and Supplemental Table S2 for ENCODE Project cell types.

For the transcription factor analysis (Fig. 1B), we downloaded processed ChIP-seq data of *POU5F1*, *SOX2*, and *NANOG* downloaded from NCBI/GEO (GSE46130). Replicate experiments were averaged. The TF peaks were overlapped with the coordinates of the differentially methylated enhancers, promoters, and islands.

For the conservation analysis of enhancer (Fig. 4B) we used 100-way PhastCons conservation data.

Copy number variations (CNVs) for skin cutaneous melanoma (SKCM) were obtained from the TCGA database as genomic segments after removal of germline CNV (TCGA\_SKCM\_GSNP6noCNV\_gSeg).

### ***Defining differentially methylated regions (DMRs)***

First, we mapped the CpG codes in the Illumina HumanMethylation450 microarray to their genomic coordinates using the microarray platform (NCBI/GEO record GPL13534; human genome release hg19/GRCh37). Second, we annotated the CpGs using the information in this record, identifying CpG sites belonging to islands, shores, shelves, enhancers and UTRs. Given that this record does not have information of CpG sites in promoters, exons or introns, we used the UCSC table 'knownGene' to identify these sites. Third, some CpG sites had several annotations, thus, we divided the CpGs into unique (non-overlapping) genomic features using the following prioritization: 1) promoters, 2) islands, 3) enhancers, 4) introns, 5) exons, 6) UTRs, 7) shores, 8) shelves and 9) intergenic regions. Fourth, for each separate genomic feature we constructed intervals using a window of 500 bp directly upstream and downstream of the CpG coordinate. Overlapping intervals (same genomic feature) were joined, and extended into a larger interval. 69% of the regions had a length of 1000 bp, 28% a length greater than 1000 bp and less than 2000 bp, no region was greater in length than 7500 bp. Fifth, we used these genomic intervals to calculate region-wise methylation levels based on the average methylation of all CpG sites within the interval, we performed this for all normal and all tumor samples in each cancer dataset. Sixth, we used the two-sample Wilcoxon test to identify differentially methylated regions (DMRs) between normal and primary samples

(methylation threshold  $>0.3$ ,  $q < 0.2$ , FDR corrected); metastatic DMRs were identified by comparing primary and metastatic melanoma (methylation threshold  $>0.2$  and  $q < 0.2$ ). The fractions of DMRs in each genomic feature presented in Fig. 1A, Fig. 3A and Supplemental Fig. S1C were normalized to the amount of intervals in each genomic feature in the Illumina HumanMethylation450 array (Dedeurwaerder et al. 2011).

### ***Gene and miRNA differential expression analysis***

A fold-change of 1.25 was used to determine differentially expressed genes and miRNAs ( $q < 0.05$ , FDR corrected; see Supplemental Methods for description).

### ***Model for prediction of gene-eDMR associations***

We developed an integrated model to predict genes or miRNAs regulated by eDMRs (Fig. 2C and flowchart in Supplemental Fig. S2H). First, we calculated the differential methylation in enhancer and promoter regions, retaining the eDMRs, and using the level of promoter methylation change in the third step. Second, we computed Spearman's correlations between eDMR methylation patterns and gene/miRNA expression patterns across the same (matched) patients. Highly correlated eDMR-genes or eDMR-miRNAs pairs were retained (negative correlation below  $-0.4$ ). Third, we filtered out genes for which promoter differential methylation was greater than 0.2, enriching for genes with expression change due to enhancer, rather than by promoter, methylation variability (performed only for predicting eDMR-associated genes and not associated-miRNAs, since annotation for miRNA promoters is incomplete). Fourth, we selected eDMR-gene and eDMR-miRNA pairs located on the same chromosome, with a maximal linear distance of 1 Mbp between the center of the enhancer and the transcription start site (for genes) or pre-miRNA start position

(for miRNAs). Fifth, we ranked eDMR-gene predictions (see Supplemental Table S4 and section *Ranking eDMR-gene pairs predictions*).

### ***Ranking eDMR-gene pairs predictions***

In order to provide confidence that the predicted eDMR-gene pairs represent active enhancer-gene interactions, we applied an integrated scoring function. Our scoring scheme combined quantitative and qualitative features of the eDMR-gene pairs. First, we ranked each eDMR-gene pair based on the following quantitative features: (a) Spearman's correlation between the eDMR and gene, (b) greater enhancer differential methylation, (c) greater gene expression fold-change, and (d) shorter eDMR-gene distances. Second, we used qualitative features which could increase ranking if existed: (a) eDMR overlap with H3K4me1, H2A.Z, or DNase I peaks, (b) promoter overlap with H3K4me3 or DNase I peaks. We summed up the scores for each eDMR-gene pairs (ties were given an average ranking score), sorted them based on their overall score, yielding ranked predictions (Supplemental Table S4).

### ***Gene enrichment analysis for diseases***

We used the DISEASES website (<http://diseases.jensenlab.org>) (Pletscher-Frankild et al. 2014) to evaluate enrichment for diseases in particular organ systems (Fig. 2E). Diseases were identified using keywords: digestive (*digest-, gastro-, gastric, stomach*), endocrine (*endocrine-, gland disease, gland cancer, gland neoplasm, thyroid, pancreas*), renal (*kidney, renal, nephron-, nephri-*), reproductive (*reproduct-, breast, prostate, uterine, cervic-, cervix, uterus*), respiratory (*lung, respirator*); duplicate genes were removed to ensure unique values for hypergeometric distribution significance testing and FDR corrected.

### ***Survival analysis***

Survival time was derived from the 'overall survival' column of the clinical data files obtained for TCGA samples. First, patients were divided into two groups (high and

low) by comparing the eDMR methylation level to the median methylation the eDMR across all patients. Similarly, for gene expression survival analysis, patients were divided into two groups by comparing to the median expression of gene across all patients. Significant differences between the two groups were determined using the Chi-squared distribution ( $q < 0.1$ , FDR corrected).

Enrichment for association of DMRs with patient survival (Fig. 3B) was determined by comparing the amount of DMRs that can significantly differentiate between patient outcomes, with the amount of static regions that can do the same (determined separately for each genomic feature).

### ***Multiple testing***

All significance tests were corrected for false discovery rates (FDR) using the Benjamini & Hochberg adjustment (Benjamini and Hochberg 1995).

### ***Computational data analysis***

Data analyses were performed using R statistical language (R Core Team 2015). We used the following packages for R in the analysis 'GenomicRanges' (version 1.16.4) (Lawrence et al. 2013), 'TxDb.Hsapiens.UCSC.hg19.knownGene' (version 3.1.2) (Carlson 2015), 'survival' (version 2.37-7) (Therneau 2015) and 'reshape2' (version 1.4.1) (Wickham 2007). Custom R scripts for determining differentially methylation regions, predicting and ranking eDMR-gene pairs are provided, together with a sample datasets derived from the TCGA are available in the Supplemental data (see 'INFO\_README.txt' file).

### ***Primary human osteoblasts and melanoma cells co culture***

Primary human osteoblasts were isolated using a protocol described by Dillon *et al.* (Dillon et al. 2012). In short, trabecular bone was obtained from healthy donors undergoing total knee arthroplasty. Written and informed consent was obtained from

all subjects. The protocol was approved by the institutional Ethics Committee at Tel-Aviv Sourasky Medical Center, in accordance with the Helsinki Declaration on the use of human subjects in research. The trabecular bone fragments were diced into small pieces and washed with sterile PBS. The diced bone extracts were then placed on a tissue culture plate with Dulbecco's Modified Eagle Medium (DMEM) supplemented with 20% fetal bovine serum, 100 U/ml penicillin, and 100 µg/ml streptomycin (all from GIBCO, Life Technologies) and incubated at 37 °C and 5% CO<sub>2</sub>. Five days later, the medium was replaced; medium was then replaced twice a week until the cells reached confluency. Osteoblasts were then seeded 24 hour prior to addition of melanoma cells in a ratio of 1:5 melanoma cells to osteoblasts). In control plates, only melanoma cells were seeded.

#### ***Cell culture and FACS sorting***

WM3682 melanoma cells were generously provided by Dr. Levi A. Garraway (Department of Medical Oncology and Center for Cancer Genome Discovery, Dana-Farber Cancer Institute, Boston, MA). Cells were cultured in DMEM medium supplemented with 10% fetal bovine serum (Sigma-Aldrich) and 1% penicillin/streptomycin/glutamine (Invitrogen). For establishment of stable cell lines, cells were transfected with GFP-expressing plasmid and selected with 1 µg/ml puromycin (Sigma-Aldrich). For co-culture, WM3682-GFP cells were cultured with bone cells for five days. Cells were collected by flow cytometry using BD FACSAria cell sorter.

#### ***RNA purification and qRT-PCR***

Total RNA was purified from sorted melanoma cells using TRIzol (Invitrogen) according to manufacturer's instructions followed by treatment with RNase-free DNase (QIAGEN). RNA was quantified based on OD<sub>260/280</sub>. For qRT-PCR analysis RNA was subjected to one-step qRT-PCR using a MultiScribe RT-PCR kit (Applied Biosystems) and FastStart Universal SYBR Green Master Mix (Roche).

### ***Plasmids and cloning***

EDMRs of the human *KIT*, *CYTL1*, and *KIF14* loci (Chr4:55708294-55709294, Chr1:201198480-201199526, Chr4:5019645-5020678, respectively) were amplified from human genomic DNA (see Supplemental Table S5 for primer sequences used in cloning). The *CYTL1* eDMR (1033 bp) and *KIF14* eDMR (1046 bp) fragments were digested with NheI-XhoI restriction enzymes and inserted into the pGL3-promoter vector (Promega) upstream of minimal promoter and firefly luciferase reporter gene (both were kindly provided by Prof. Eran Bacharach, Department of Cell Research and Immunology, Faculty of Life Sciences, Tel-Aviv University, Israel). The *KIT* eDMR (1000 bp) fragment was cloned into PGL3-promoter reporter plasmid digested with SmaI-XhoI restriction enzymes. pCDNA3-*KIT* expression vector was kindly obtained from Prof. Lars Rönnstrand (Division of Translational Cancer Research and Lund Stem Cell Center, Lund University, Medicon Village, SE-22381 Lund, Sweden).

### ***In vitro methylation, transfections and dual luciferase assay***

The *KIT*, *CYTL1*, and *KIF14* eDMR firefly luciferase reporter vectors were *in vitro* methylated using the methylase SssI (New England Biolabs), according to manufacturer's recommendations, followed by purification using Wizard SV PCR clean-up system (Promega). Successful methylation was verified by restriction enzyme digestion with the methylation-sensitive (HpaII) and methylation-insensitive (MspI) enzymes and (New England Biolabs). The digestion patterns were analyzed by agarose gel electrophoresis. WM3682 melanoma cell lines were co-transfected using jetPEI™, with methylated eDMR- luciferase reporter plasmid, unmethylated reporter, or reporter plasmid without eDMR (as control) and with pRL-plasmid (Promega). Luciferase activity was measured 48 hours after transfection using Dual Luciferase kit (Promega). Firefly luciferase activity was normalized to the *Renilla* luciferase.

### ***Invasion assay***

WM3682 melanoma cell lines were transfected with *KIT* expression vector or empty vector (as control) using jetPEI™. 48 hours post transfection invasion assay was performed as previously described (Golan et al. 2015).

### ***5-aza-deoxycytidine treatment***

WM3682 melanoma cell lines were treated with 10 μM 5-aza-dC (Sigma-Aldrich) for 48 hours, following by RNA purification and qRT-PCR as described. Baseline expression was established by mock treatment of cells with DMSO.

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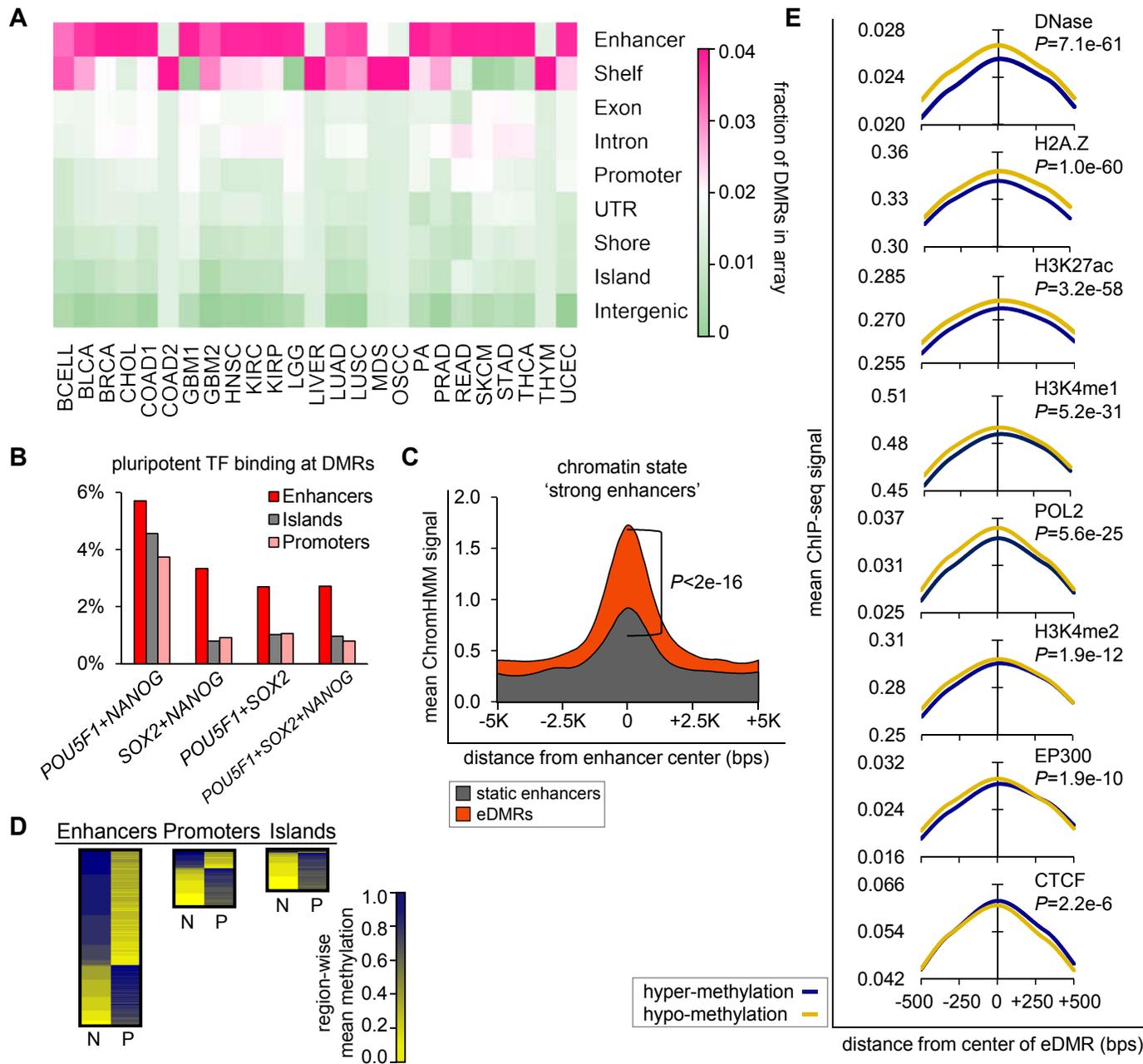


Figure 1: Enhancers exhibit highly dynamic methylation patterns upon malignant transformation

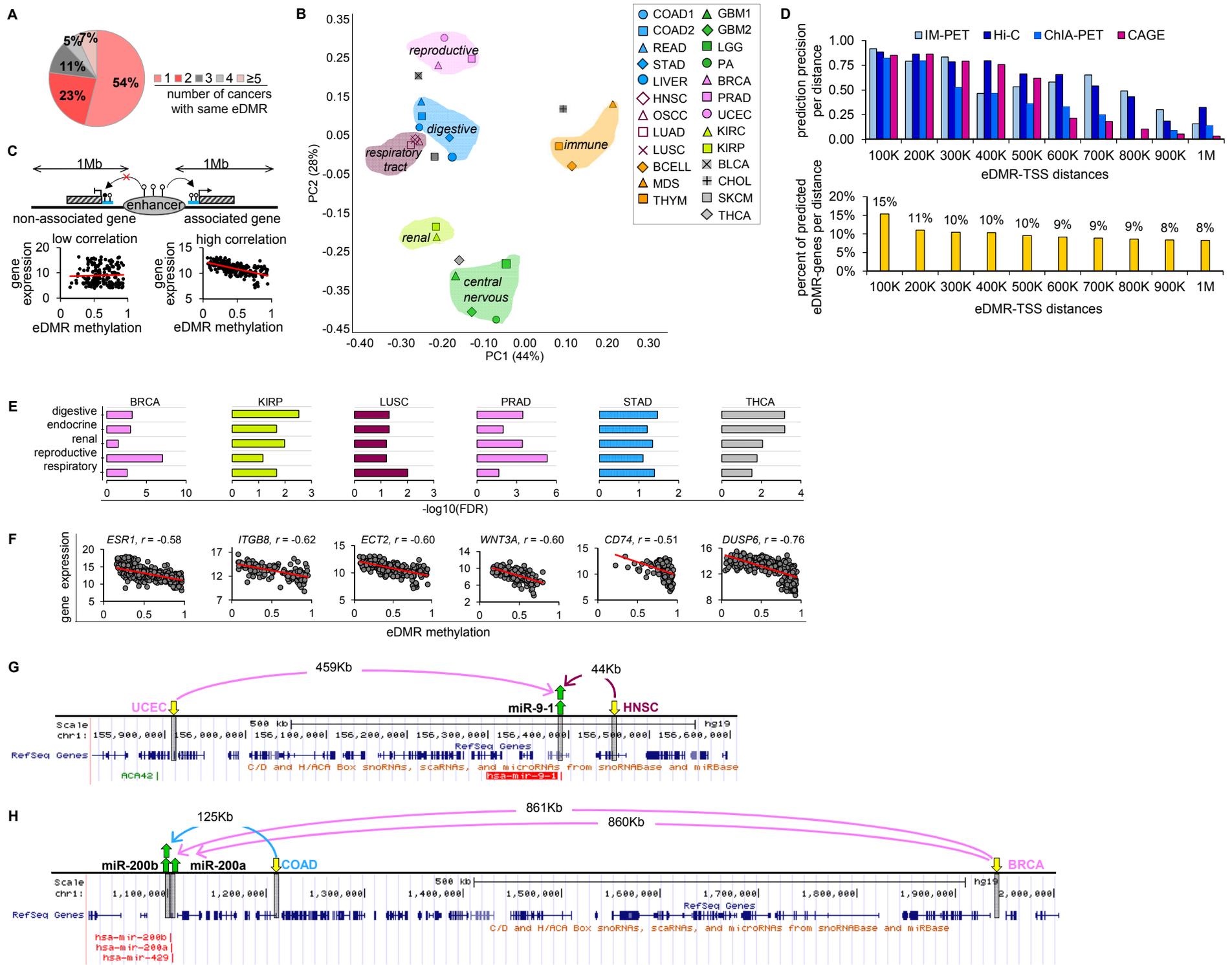


Figure 2: Variation in enhancer methylation classifies primary tumors according to their organ systems

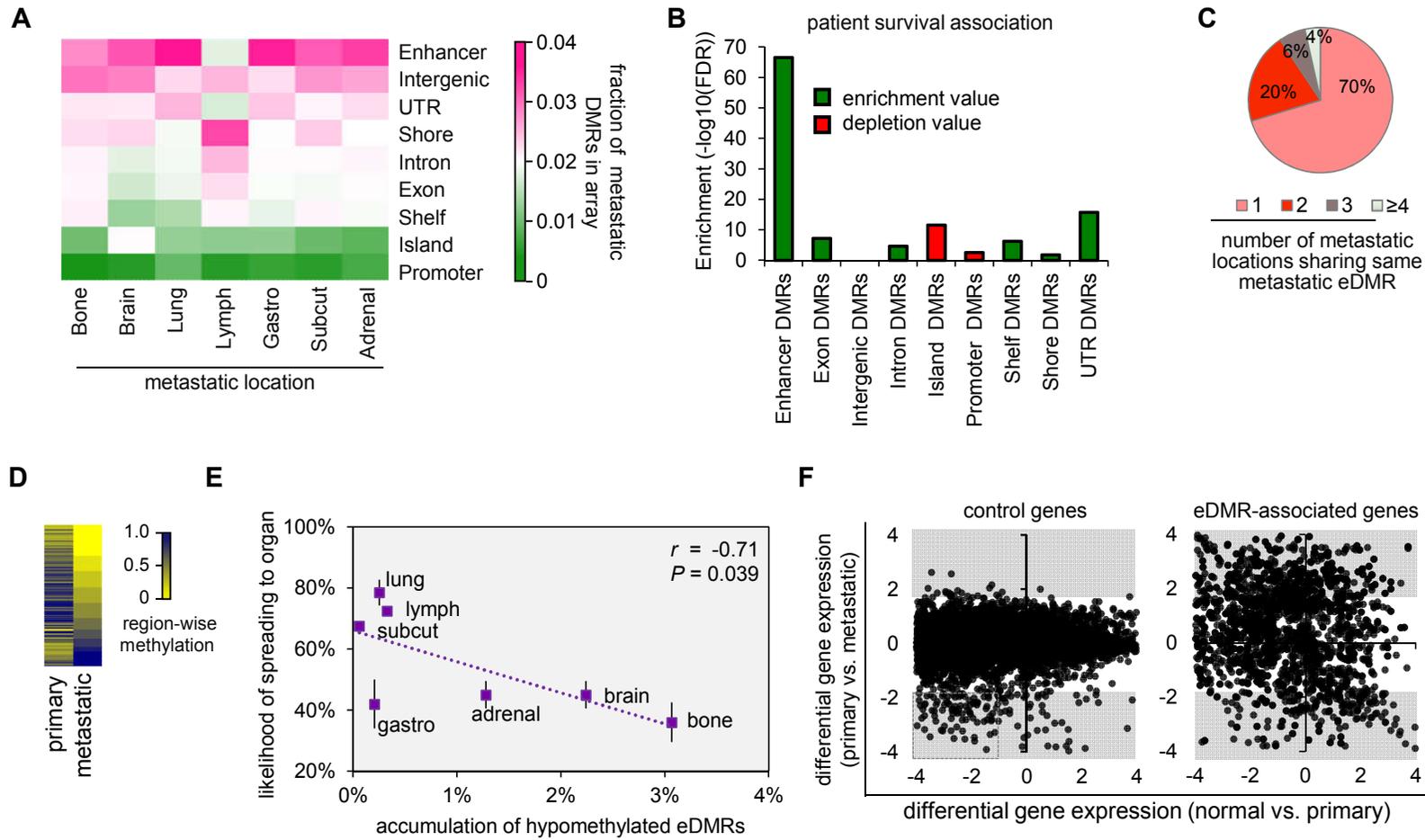


Figure 3: Accumulation of eDMR hypomethylation correlates with likelihood of metastasis and patient outcome

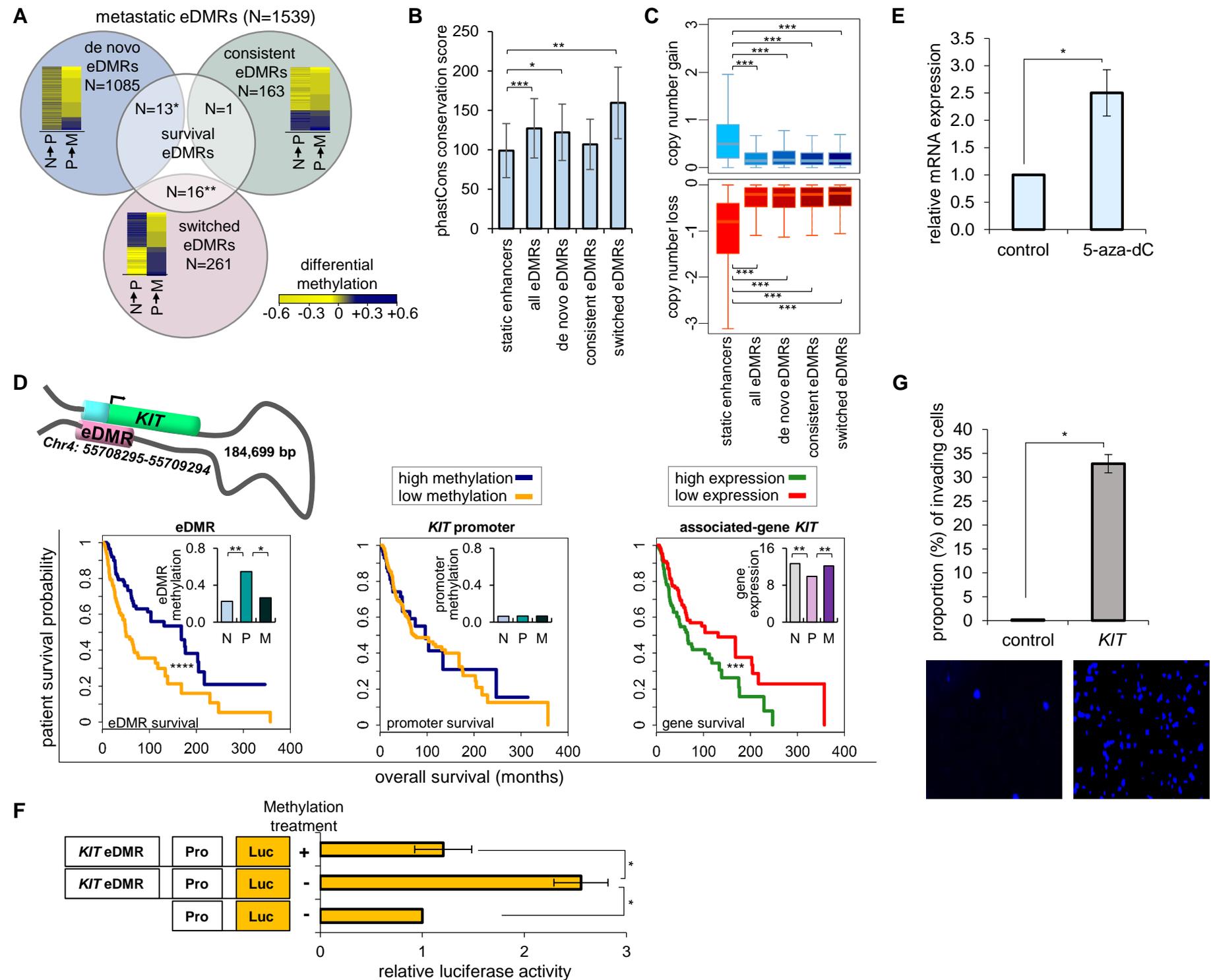


Figure 4: Methylation plasticity of melanoma eDMRs is associated with patient mortality

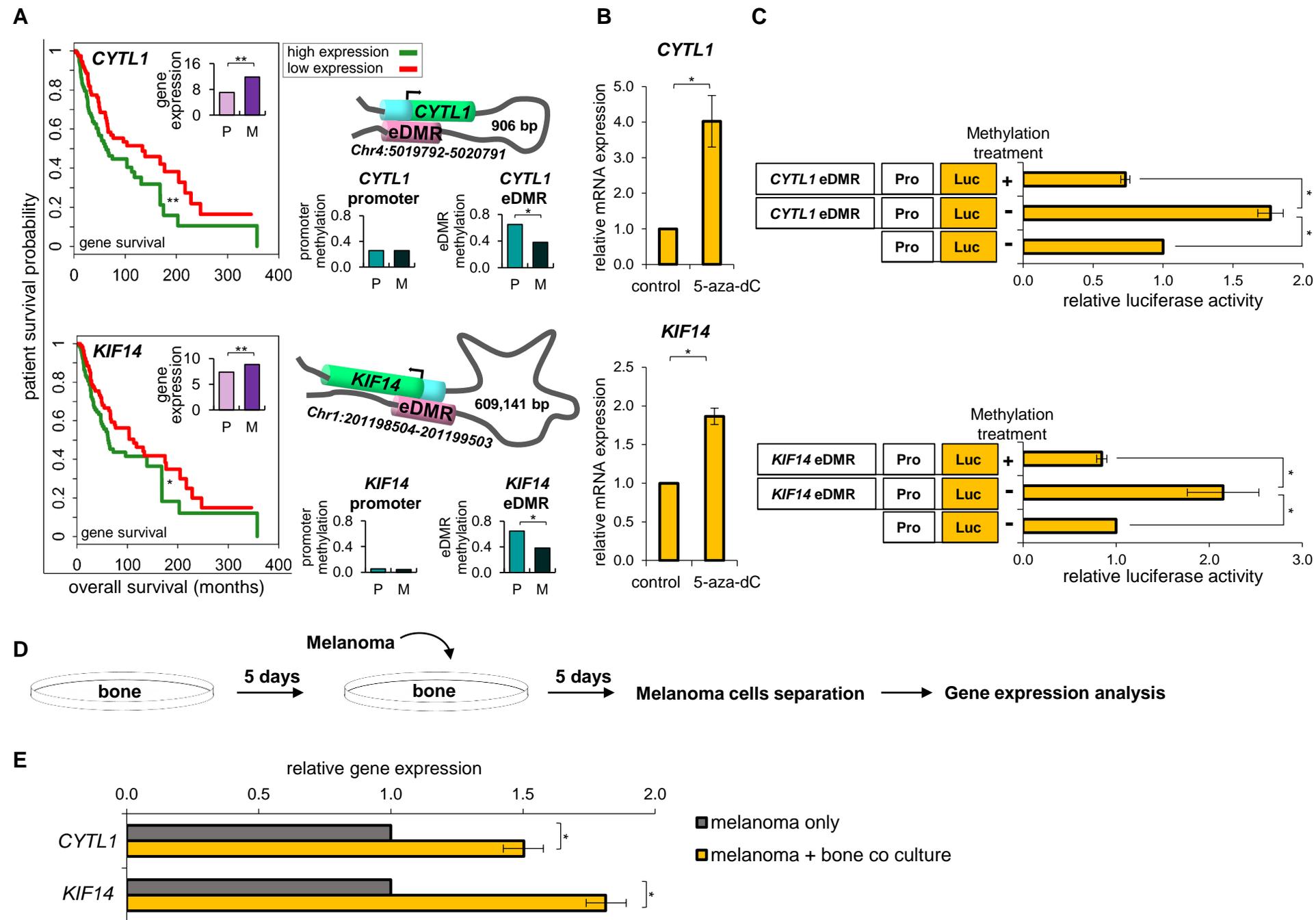


Figure 5: Melanoma co-culture with osteoblasts alters pro-cancer eDMR-associated gene expression



## Enhancer methylation dynamics contribute to cancer plasticity and patient mortality

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