The Effect of CDK6 Expression on DNA Methylation and DNMT3B Regulation

Cdk6 is bound to the 5’ region of Dnmt3b

ChIP-seq

Cdk6

TF

Dnmt3b

Cdk6 loss affects Dnmt3b expression ...

RNA-seq

Chromatome

Cdk6

TF

Dnmt3b ↓

... and DNA methylation

RRBS

Cdk6

methylated

unmethylated

HIGHLIGHTS

- CDK6 and DNMT/TET/HDAC expression correlate in ALL samples
- CDK6 transcriptionally regulates Dnmt3B in BCR-ABL+ cells
- DNA methylation patterns change upon CDK6 loss in BCR-ABL+ cells
The Effect of CDK6 Expression on DNA Methylation and DNMT3B Regulation

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SUMMARY
CDK6 is frequently overexpressed in various cancer types and functions as a positive regulator of the cell cycle and as a coregulator of gene transcription. We provide evidence that CDK6 is involved in the process of DNA methylation, at least in ALL. We observe a positive correlation of CDK6 and DNMT expression in a large number of ALL samples. ChIP-seq analysis reveals CDK6 binding to genomic regions associated with DNA methyltransferases (DNMTs). ATAC-seq shows a strong reduction in chromatin accessibility for DNMT3B in CDK6-deficient BCR-ABL+ Cdk6−/− cells, accompanied by lower levels of DNMT3B mRNA and less chromatin-bound DNMT3B, as shown by RNA-seq and chromatome analysis. Motif analysis suggests that ETS family members interact with CDK6 to regulate DNMT3B. Reduced representation bisulfite sequencing analysis uncovers reversible and cell line-specific changes in DNA methylation patterns upon CDK6 loss. The results reveal a function of CDK6 as a regulator of DNA methylation in transformed cells.

INTRODUCTION
The transformation of normal to cancer cells is driven by a multitude of genetic and epigenetic alterations. There is recent evidence that overexpression of the cyclin-dependent kinase 6 (CDK6) is a frequent event in hematological malignancies and solid tumors (Tadesse et al., 2015; Nagel et al., 2008; Tigan et al., 2016). Functional characterization of CDK6 has revealed that it affects a variety of molecular processes (Bellutti et al., 2018; Scheicher et al., 2015; Uras et al., 2017, 2019; Kollmann et al., 2013; Wang et al., 2017). Besides binding D-type cyclins to drive the G1 phase of the cell cycle, it also regulates gene expression (Kollmann et al., 2013; Tigan et al., 2016). Independent of its kinase activity, it interacts with transcription factors to act as a chromatin-bound cofactor that induces or represses the expression of specific genes. We have shown that CDK6 interacts with STAT3 to induce the expression of p16INK4a and with the AP-1 transcription factor c-Jun to induce the expression of VEGF-A (Kollmann et al., 2013). It also cooperates with the NF-κB subunit p65 to induce an inflammatory gene response (Handschick et al., 2014). CDK6 can also repress transcription, as it does in hematopoietic stem cells by interacting with AP-1 to repress EGR1 expression and thereby to induce the exit from quiescence and the growth of leukemic stem cells (Scheicher et al., 2015). Together with NFYA and SP1, CDK6 induces a transcriptional program to block p53 in hematopoietic cells (Bellutti et al., 2018).

Gene transcription is frequently regulated by epigenetic mechanisms, such as DNA methylation, various chemical modifications of histone proteins and chromatin restructuring, which results in gene activation or gene silencing (Sandoval and Esteller, 2012). The main targets of methylation in the mammalian genome are cytosines within CpG dinucleotides. CpG dinucleotides are distributed throughout the genome but are concentrated in regions of about 0.5–2 kb in length, called CpG islands, that are associated with ~60% of human gene promoters (Sandoval and Esteller, 2012). DNA methylation is considered important in the pathogenesis of many solid tumors and hematological malignancies (Heller et al., 2016; Guillamot et al., 2016; Kimura et al., 2019). The CpG islands of many cancer-associated genes are frequently methylated in cancer cells, resulting in the transcriptional silencing of these genes (Sandoval and Esteller, 2012; Heller et al., 2013). Changes in DNA methylation in regions outside CpG islands may be equally important in leukemogenesis, with hypomethylation as relevant as hypermethylation (Guillamot et al., 2016; Figueroa...
Despite a wealth of evidence, the mechanisms of aberrant methylation in cancer cells remain unclear. There is a preliminary indication that CDK6 may play a part, as inhibition of CDK kinase with palbociclib promotes the degradation of the DNA methyltransferase 1 (DNMT1) (Acevedo et al., 2016). We now confirm that CDK6 is involved in the regulation of DNA methylation, although its function does not depend on its kinase activity. In several independent gene expression microarray datasets from a large number of ALL samples CDK6 is co-expressed with DNMTs, TET1 and histone deacetylases (HDAC). ChIP-seq, ATAC-seq, RNA-seq, and mass spectrometry analysis of chromatin (referred to as chromatome) reveal that DNMT3B is a target for transcriptional regulation by CDK6 in BCR-ABL+ cells. Reduced representation bisulfite sequencing (RRBS) shows that loss of CDK6 causes changes in DNA methylation patterns in a cell line-specific manner, with both hypomethylation and hypermethylation prominent. Many of the changes in methylation can be reversed by re-expressing CDK6. The findings demonstrate that CDK6 has a function as a regulator of DNA methylation in transformed cells.

**RESULTS**

**CDK6 and DNMT/TET/HDAC Expression Correlate in ALL Samples**

Epigenetic changes are frequent in leukemia and epigenetic regulators such as DNMTs and TETs are often mutated in hematopoietic malignancies (Yang et al., 2015, Delhommeau et al., 2009, Cancer Genome Atlas Research Network et al., 2013, Kimura et al., 2019, de Keersmaecker et al., 2013). Coexpression analysis using four independent gene expression microarray data sets from ALL patients showed a positive correlation between levels of CDK6 and of DNMT3A (mean R = 0.46 in 4 data sets; p < 0.0001), with an only slightly weaker positive correlation with levels of DNMT3B (mean R = 0.42 in 3 of 4 data sets; p < 0.0001) and a weaker correlation with levels of DNMT1 (mean R = 0.34 in 4 data sets; p < 0.0001; Figure 1). While CDK6 was positively correlated with TET1 (mean R = 0.49 in 4 data sets; p < 0.0001) and negatively with TET2 (mean R = −0.35 in 4 data sets; p < 0.0001), there was no correlation with TET3 in ALL patients (Figure 1).

HDACs are also affected in hematopoietic malignancies, although aberrant expression is more common than mutations (Moreno et al., 2010; Zhang et al., 2015). Of the large family of HDACs, we found HDAC1 and HDAC8 to be coexpressed with CDK6 in ALL patients (mean R = 0.55 and 0.42 in 4 data sets; p < 0.0001, respectively). The patterns of coexpression between CDK6 and DNMT/TET/HDACs are highly similar in the four gene expression data sets (Figure 1), showing that expression of CDK6 and several factors associated with DNA methylation are positively correlated in ALL. The data suggest that CDK6 might be involved in the changes in DNA methylation during leukemogenesis.

**DNMT3B Is Transcriptionally Regulated by CDK6 in BCR-ABL+ Cells**

The transcriptional regulator CDK6 is particularly significant in leukemia, where it is upregulated and contributes to leukemogenesis (Nagel et al., 2008; Tadesse et al., 2015; Tigan et al., 2016). The positive correlation between expression of CDK6 and DNMT3 prompted us to study whether DNMT1, DNMT3A and DNMT3B are transcriptionally regulated by CDK6. Murine (C57Bl/6J) Cdk6wt and Cdk6−/− cell lines obtained from single-cell bone marrow suspension were retrovirally transduced with a pMSCV-BCR-ABLp185-IRE5-GFP vector as described (Belluti et al., 2018). CDK6 ChIP-seq analysis of these cells revealed specific ChIP peaks in the 5′ regions of DNMT1 and DNMT3B (Figures 2A and S1A) and additional CDK6 peaks in the promoter and exonic/intronic regions of DNMT3A (Figure S1B). The chromatin accessibility of the genomic regions harboring the DNMT1 and DNMT3B genes also depended on the presence of CDK6. ATAC-seq analysis in BCR-ABL+ Cdk6wt and BCR-ABL− Cdk6−/− cells showed that the chromatin accessibility in the DNMT3A region was comparable. However, the chromatin accessibility in the DNMT1 and DNMT3B regions was strongly reduced in CDK6-deficient BCR-ABL+ Cdk6−/− cells (Figures 2A and S1A). The significance of the differences in ATAC-seq were confirmed by RNA-seq analysis, which revealed reduced DNMT3B expression in BCR-ABL+ Cdk6−/− cells (Figure 2B), while expression of DNMT1 and DNMT3A remained unchanged.

Analysis of the chromatin-bound proteome showed that the changes in transcription are reflected by alterations in protein abundance: less DNMT3B bound to chromatin in BCR-ABL− Cdk6−/− cells than in BCR-ABL+ Cdk6wt cells (Figure 2C). Motif analysis of the genomic region underlying the CDK6
ChIP-seq/ATAC-seq peak (Figure 2A) identified ETS motifs overlapping with CDK6 binding sites in the 5' region of DNMT3B (Figure 2D). To test if ETS1 binds to this genomic location, we analyzed ETS1 ChIP-seq data from human K562 cells and murine B-cells. We verified that an ETS1 peak overlaps with the CDK6 binding site in the promoter/exon 1 region of DNMT3B in both data sets (Figure 2E). These findings suggest that CDK6 directly regulates DNMT3B, possibly mediated by ETS factors.

DNA Methylation Patterns Change upon Loss of CDK6 in BCR-ABL+ Cells

To investigate whether the alterations in DNMT3 expression provoke changes of the methylome, we used a CRISPR-Cas9-based strategy to generate BCR-ABL+ cells deficient for CDK6 as confirmed by Western blotting (Figure S2). Our procedure (for details see Transparent Methods) maintains genetic stability and avoids altering p53: CDK6-deficient transformed cells consistently harbor p53 mutations that may affect the methylome (Tovy et al., 2017). Dnmt3b expression was downregulated upon Cdk6 loss in these cells (Figures 3A and S3). Further, Cdk6 reexpression strongly induced Dnmt3b mRNA expression (Figure 3A). We used 3 individually derived BCR-ABL+ cell lines for RRBS to study the methylome. Principal component analysis (PCA) based on CpG site methylation unequivocally separated BCR-ABL+Cdk6+/C0/C0 cells from BCR-ABL+Cdk6wt cells (Figure 3B). The separation was also evident in the sample correlation plot, which showed weaker correlation between BCR-ABL+Cdk6+/C0/C0 and BCR-ABL+Cdk6wt cells than within BCR-ABL+Cdk6wt cells (Figure S4).

Unexpectedly, the BCR-ABL+Cdk6+/C0/C0 samples are not represented in a distinct cluster in the PCA but are separated from one another, indicating that loss of CDK6 is accompanied by different changes in DNA
methylations in the different cell lines. To investigate the differences in more detail, we examined differential methylation between individual pairs of cell lines. We found 4,707 (2,028 hyper, 2,679 hypo), 4,565 (2,498 hyper, 2,067 hypo) and 5,528 (2,773 hyper, 2,756 hypo) CpG sites with a difference in methylation of at least 40% in each of the three pairwise comparisons (Figure 3C). Only ~1.5% of differentially methylated CpG

Figure 2. DNMT3B Is Transcriptionally Regulated by CDK6 in BCR-ABL⁺ Cells
(A) Binding of CDK6 to the 5' region of DNMT3B in BCR-ABL⁺ cells determined by ChIP-seq analysis (upper panel). Chromatin accessibility at the 5' region of DNMT3B in BCR-ABL⁺ Cdk6⁺⁺ (green) and BCR-ABL⁺ Cdk6⁻⁻ (red) cells determined by ATAC-seq analysis. See also Figure S1.
(B) Heatmap showing mRNA expression of DNMT1/3a/3b in BCR-ABL⁺ Cdk6⁺⁺ (green, N = 3) and BCR-ABL⁺ Cdk6⁻⁻ (red, N = 3) cells analyzed by RNA-sequencing.
(C) Chromatin-bound DNMT1/3a/3b in BCR-ABL⁺ Cdk6⁺⁺ (green) and BCR-ABL⁺ Cdk6⁻⁻ (red) cells analyzed by chromatome analysis. Mean values + SD are shown.
(D) ETS motifs found within the ATAC-seq peak at the 5' UTR of DNMT3B.
(E) Binding of ETS1 to the 5' region of DNMT3B in human K562 cells (left panel) and in murine spleen cells (right panel) determined by ChIP-seq analysis. See also Figure S1.
Figure 3. Analysis of the Methyome of BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{wt} and BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{-/-} Cells By Reduced Representation Bisulfite Sequencing

(A) Quantification of Dnmt3b mRNA expression in BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{wt}, BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{-/-} and BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{-/-}+Cdk6 cells determined by RT-PCR. Mean fold changes +SEM are shown (N = 8), p values were calculated using one-sample t-tests.

(B) Principal component analysis of 3 BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{wt} (WT, red) and 3 BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{-/-} (KO, blue) cell lines based on CpG methylation. Each dot represents a unique sample. See also Figure S4.

(C) Volcano plots showing hypomethylated and hypermethylated CpG sites in 3 replicates of BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{-/-} cell lines compared to BCR-ABL\textsuperscript{+} Cdk6\textsuperscript{wt} cells.
Figure 3. Continued

(D) Venn diagram illustrating the overlap of differentially methylated CpG sites in the 3 BCR-ABL+ Cdk6−/− cell lines.
(E) Density plot showing the location of hypermethylated (pink) and hypomethylated (blue) CpG sites relative to transcriptional start sites (TSS). See also Figure S4.

sites were shared in two of the cell lines and no single CpG site showed a consistent difference in methylation in all 3 cell lines (Figure 3D). Annotation of the differentially methylated CpG sites to genomic region revealed that both hyper- and hypomethylated CpG sites are closely associated with transcriptional start sites (TSSs) of genes (Figure 3E). The data confirm that loss of CDK6 has a dramatic effect on DNA methylation in BCR-ABL+ cells and suggest that the precise nature of the effect depends on stochastic processes.

In the next step, we asked if DNA methylation changes are also associated with CDK6 expression in patients. Thus, we stratified the TCGA LAML data set into CDK6-low (lower quartile, N = 27) and CDK6-high (upper quartile, N = 27) samples and tested for correlation between CDK6 and DNMT3B expression as well as for differences in DNA methylation. As for ALL samples (Figure 1), we found a positive correlation between CDK6 and DNMT3B expression in AML samples (R = 0.47, p < 0.005). Further, CDK6-high samples showed a strong increase in methylation (β-difference >0.3) for 1,287 probes (Figure S5). Only 11 probes with decreased methylation (β-difference < -0.3) were found indicating that CDK6 expression is mainly associated with hypermethylation in patient samples.

Expression of CDK6 Reverses Changes in CpG Methylation Caused by Loss of CDK6

It is conceivable that the variable changes in DNA methylation are a result of different levels of CDK6. We tested this possibility by re-expressing CDK6 using retroviral transduction with a construct encoding HA-CDK6 (Figure S2) and RRBS analysis and compared methylation in the reconstituted BCR-ABL+ Cdk6−/−+ Cdk6 cells and the maternal cells (BCR-ABL+ Cdk6wt and BCR-ABL+ Cdk6−/−). Re-expression of CDK6 did not result in the formation of a separate methylation cluster in the hierarchical cluster analysis but showed that BCR-ABL+ Cdk6−/−+ Cdk6 cells clustered with BCR-ABL+ Cdk6−/− cells from the same background (Figure 4A). Thus, re-expression of CDK6 does not induce global changes of the methylome but regulates a distinct set of genes in each cell line. We then analyzed whether the methylation changes caused by loss of CDK6 are reverted by re-expression of CDK6. Hypermethylation of 79% (range: 70%–85%) of CpG sites was found to be reversible (>1.5-fold increase in methylation) in the three cell lines, while methylation of 34% of the hypermethylated CpG sites (mean: 34%; range: 26%–48%) was reverted when CDK6 is re-expressed (Figure 4B). The reversal of methylation patterns was independent of the location of CpG sites and was found consistently in CGIs, CGI shores, CGI shelves, inter CGI regions, promoters, introns, exons, and intergenic regions (Figure S6). The effect of re-expression of CDK6 on the methylation of 5,000 randomly selected CpG sites was weak (mean methylation changes: 6.6%; repeated 10 times), confirming that the changes in methylation mediated by CDK6 re-expression are specific (Figure S7).

As the re-expression of CDK6 produces varying amounts of the protein in the individual cells, the consistent nature of the changes in methylation caused by re-expression of CDK6 shows that the changes are independent of the level of CDK6. In other words, expression of CDK6 is associated with specific changes in CpG methylation, with the precise effects different for each individual clone. Re-expression of CDK6 largely reverts the hypomethylation mediated by loss of CDK6, although the hypermethylation in BCR-ABL+ Cdk6−/− cells is only partially reversible, strengthening the dominating hypermethylation seen in CDK6-high patient samples (Figure S5).

CDK6-Dependent Differential Methylation Is Associated with Altered Gene Expression in BCR-ABL+ Cells

Loss of CDK6 is thus associated with changes in DNA methylation. As not all changes in DNA methylation result in differences in gene regulation, we performed RNA-seq analysis to examine the consequences of loss of CDK6 on gene expression. PCA based on all gene expression data revealed that BCR-ABL+ Cdk6−/− cells and BCR-ABL+ Cdk6wt cells cluster separately with lower heterogeneity than the RRBS clustering (Figure S8). The overlay of CDK6-dependent methylation and transcriptional regulation uncovered an inverse pattern of decreased methylation/increased gene expression or increased methylation/decreased gene expression for 12 (2 up, 10 down), 29 (16 up, 13 down) and 14 (8 up, 6 down) genes in the three cell lines (Figure 5A). In silico functional characterization of these genes identified them as involved in cell differentiation (e.g. Runx2, Shox2, Shb), cell proliferation (e.g. Fzd3, Satb1, TBX2), immune systems (e.g. Gfi1,
NFκB, MAPK signaling (e.g., Dusp4, Mlkl), Wnt signaling (e.g., Fzd3, Fzd6) or cell growth regulation (e.g., Lhx2, Bcl6) (Figure 5B). This shows that various genes involved in cancer-related pathways are transcriptionally deregulated by CDK6-mediated changes in DNA methylation.

**DISCUSSION**

We show for the first time that CDK6 directly affects DNA methylation in a cell-specific manner. Global demethylation and de novo methylation of selected CpG islands are key events in the pathogenesis of malignant diseases (Michalak et al., 2019; Zhou et al., 2018). During the past decade, tremendous effort has been spent to identify the mechanisms of these epigenetic alterations in cancer cells and a variety of molecular changes have been associated with deregulated DNA methylation. Large-scale studies have shown that certain genes encoding enzymes involved in DNA methylation are affected by somatic mutations in certain types of cancer (Cancer Genome Atlas Research Network et al., 2013; Walter et al., 2011; Ley et al., 2010). Mutations of DNA methyltransferases (DNMTs), especially of DNMT3A, TET2, IDH1, and IDH2, have been found in AML and other hematopoietic malignancies (Yang et al., 2015; Spencer et al., 2017; Delhommeau et al., 2009; Paschka et al., 2010, Cancer Genome Atlas Research Network et al., 2013) and deregulated expression of DNMTs has been found in multiple tumor types, resulting in aberrant DNA methylation (Esteller, 2008; Peng et al., 2006; Saito et al., 2003).

We show that expression of DNMT3 and other epigenetic modifiers correlates with CDK6 expression in ALL patients. To investigate the mechanism we used a multi-omics approach in BCR-ABL+ cells. ChIP-seq
analysis showed that CDK6 is located at multiple sites within the 5’ regions and/or intragenic regions of DNMTs. In the case of DNMT3B, ATAC-seq and RNA-seq analysis associated CDK6 binding with open chromatin and transcriptional regulation. The regulation of DNMT3B mediated by CDK6 is reflected on the protein level and chromatome analysis showed reduced levels of DNMT3B protein in CDK6-deficient cells. Of note, we observed a correlation between CDK6 and TET1 expression in the ALL datasets. However, because TET1 expression was not detected in our experimental system, this correlation was not further investigated.

While the frequency of DNMT3B mutations in cancer cells is generally low, deregulated expression of this gene is of potential clinical relevance in multiple cancer types (Hayette et al., 2012; Lamba et al., 2018;
Overexpression of DNMT3B has been correlated with shorter event-free survival (EFS) and a trend toward poorer overall survival (OS) in a large panel of de novo AML patients (Hayette et al., 2012). Increased expression of DNMT3B has also been associated with poor clinical outcome, worse minimal residual disease, high rate of relapse or resistant disease and worse EFS in pediatric AML patients (Lamba et al., 2018) and older adults with primary, cytogenetically normal AML, and high levels of DNMT3B had fewer complete remissions, inferior disease-free survival, and shorter OS (Niederwieser et al., 2015). Prognostically relevant overexpression of DNMT3B is not confined to AML patients. It has also been described for patients with diffuse large B-cell lymphomas and DNMT3B overexpression was detected in clinical specimens from T-ALL and Burkitt's lymphoma patients (Poole et al., 2017; Amara et al., 2010). We observed heterogeneous expression of DNMT3A/B, with levels positively correlating with levels of CDK6 in adult or pediatric ALL samples (Figure 1). This supports the idea that the regulatory effect of CDK6 is not confined to our in vitro experiments but is also relevant in vivo.

CDK6 does not have a DNA-binding domain, so can only bind to DNA in conjunction with transcription factors. Such cooperations have been observed between CDK6 and STAT3, c-Jun, p65, AP-1, NFYA, and SP1 and result in the transcriptional regulation of specific target genes (Kollmann et al., 2013; Handschick et al., 2014; Scheicher et al., 2015; Bellutti et al., 2018). To determine which transcription factor interacts with CDK6 on the 5’ region of DNMT3B, we performed motif analysis of the region underlying the ChIP-seq/ATAC-seq peak as shown in Figure 2A. ETS motifs are the predominant binding sites and ETS1 ChIP-seq analysis identified ETS1 peaks in the DNMT3B 5’ region in human BCR-ABL+ K562 cells as well as murine B-cells. These data suggest that DNMT3B may be regulated by the binding of CDK6/ETS family to its promoter region. This idea will be addressed in further studies.

To determine whether CDK6 affects the methylome we used the next-generation sequencing approach RRBS to perform extensive methylation analysis in BCR-ABL+ cells either expressing or lacking CDK6. We found both hypermethylation and hypomethylation. Although the numbers of differentially methylated CpG sites in different cell lines were comparable, we failed to detect a common pattern of differential methylation. It is attractive to speculate that this observation reflects the heterogeneity in tumor DNA methylation that has been described for various types of human cancer (Wenger et al., 2019; Sheffield et al., 2017; Brocks et al., 2014; Li et al., 2016; Landau et al., 2014; Nordlund et al., 2013). Our result is consistent with the report that aberrant DNA methylation is a signature for the heterogeneity of ALL patients of similar cytogenetic backgrounds (Nordlund et al., 2013). Importantly, many of the methylation changes mediated by CDK6 can be reverted by re-expressing CDK6. This indicates that CDK6 is directly involved in the process of DNA methylation rather than inducing secondary methylation changes. Our finding is consistent with indications that CDK6 is involved in epigenetic programming, such as the observation that DNMT1 is degraded by inhibition of CDK6 with the CDK4/6 inhibitor palbociclib (Acevedo et al., 2016) and that the CDK4/6 inhibitor abemaciclib causes cell cycle inhibition and changes DNA demethylation and immunogenicity (Dowless et al., 2018). However, we provide the first evidence of a direct effect of CDK6 on methylation.

The molecular consequences of the changes in DNA methylation mediated by CDK6 are unclear. It is possible that CDK6 does more than regulate gene transcription, conceivably also having effects on transcription factor binding and chromatin organization. DNA methylation can either mask consensus DNA-binding sites or create new binding sites for transcription factors and nuclear complexes (Bartke et al., 2010; Spruijt et al., 2013). The transcription factor CTCF, which is involved in large-scale chromatin organization, is known to be methylation-sensitive (Bell and Felsenfeld, 2000; Hashimoto et al., 2017; Ghirlanda and Felsenfeld, 2016). DNA methylation prevents CTCF binding, thus, CDK6-mediated changes in DNA methylation may be associated with establishing and/or maintaining higher-order topological structures in the genome.

In summary, we show that loss of CDK6 contributes to DNA hypomethylation and hypermethylation. Importantly, the effect of CDK6 on DNA methylation is generally reversible, a point that should be considered when targeting CDK6 in anticancer therapy. We also identify DNMT3B as a transcriptional target of CDK6. This function of CDK6 was unknown so far but the extent to which CDK6-dependent changes in DNA methylation contribute to tumorigenesis and the maintenance of a malignant phenotype remains to be determined.
Limitations of the Study

In the present study, we demonstrated that CDK6 affects DNA methylation by transcriptional regulation of DNMT3B. However, because CDK6 does not have a DNA-binding domain, it can only bind to DNA in association with transcription factors. Our data suggest that ETS1 is one of these factors but the interaction between CDK6 and ETS1 remains to be investigated in more detail.

Resource Availability

Lead Contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Veronika Sexl (Veronika.Sexl@vetmeduni.ac.at).

Materials Availability

This study did not generate new unique reagents.

Data Availability and Code Availability

The accession numbers for the data sets reported in this paper are: GEO: GSE145220, GEO: GSE156966, GEO: GSE145220, GEO: GSE113752.

METHODS

All methods can be found in the accompanying Transparent Methods supplemental file.

SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at https://doi.org/10.1016/j.isci.2020.101602.

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AUTHOR CONTRIBUTIONS

GH performed bioinformatic data analysis, data visualization, and wrote the manuscript; SN, FB, HÜ, LS, and KK performed wet lab experiments; MZ performed bioinformatic data analysis; VS and KK supervised the project team and wrote the manuscript; all authors carefully proof-read the manuscript.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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Supplemental Information

The Effect of CDK6 Expression
on DNA Methylation
and DNMT3B Regulation

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Figure S1, related to Figure 2. Binding of Cdk6 to the 5’ regions of (A) Dnmt1 and (B) Dnmt3a in BCR-ABL+ cells determined by ChIP-seq analyses (upper panel). Chromatin accessibility at the 5’ regions of Dnmt1 and Dnmt3a in BCR-ABL+ Cdk6wt (green) and BCR-ABL+ Cdk6−/− (red) cells determined by ATAC-seq analyses (lower panel).
Figure S2, related to Figure 3. Western blot analyses to confirm Cdk6 loss and Cdk6 re-expression upon Cdk6 knock-out and HA-Cdk6 transduction, respectively. Results from 3 different cell lines (Rep 1-3) are shown. kD, kilodalton; M, marker.
Figure S3, related to Figure 3. Western blot analyses of Dnmt3b and Cdk6 in BCR-ABL+ Cdk6+/+ and BCR-ABL+ Cdk6−/− cell lines. Signals were densitometrically quantified and normalized to HSC70. kD, kilodalton; M, marker.
Figure S4, related to Figure 3. Correlation plot of 3 BCR-ABL+ Cdk6wt (WT) and 3 BCR-ABL+ Cdk6−/− (KO) cell lines based on CpG methylation.
Figure S5, related to Figure 3. Heatmap showing methylation of 1,298 probes with a β-difference of at least +/-0.3 between CDK6\textsuperscript{high} (upper quartile, N = 27) and CDK6\textsuperscript{low} (lower quartile, N = 27) AML samples from the TCGA LAML dataset. Values range from 0 (unmethylated; blue) to 1 (fully methylated; red). Each row represents a unique probe on the Infinium Human Methylation 450K BeadChip and each column represents a unique patient.
Figure S6, related to Figure 4. Heatmaps showing methylation values of differentially methylated CpG sites in 3 replicates (Rep 1-3) of BCR-ABL Cdk6<sup>wt</sup>, BCR-ABL Cdk6<sup>−/−</sup> and BCR-ABL Cdk6<sup>−/−</sup>+Cdk6 cell lines separated for CpG islands (CGI), CGI shores, CGI shelves, inter CGI regions, promoter regions, exons, introns and intergenic regions. Values are depicted as percentage of methylation and range from 0% (dark blue) to 100% (dark red).
Figure S7, related to Figure 4. Heatmaps showing methylation values of 5,000 randomly selected CpG sites in BCR-ABL<sup>+</sup> Cdk<sup>6<sup>wt</sup></sup>, BCR-ABL<sup>+</sup> Cdk6<sup>−/−</sup> and BCR-ABL<sup>+</sup> Cdk6<sup>−/−</sup>+Cdk6 cell lines. Random CpG site selection was repeated for 10 times. Values are depicted as percentage of methylation and range from 0% (dark blue) to 100% (dark red).
Figure S8, related to Figure 5. Principal component (PC) analysis of 3 BCR-ABL⁺ Cdk6⁺/⁻ (WT, red) and 3 BCR-ABL⁺ Cdk6⁻/⁻ (KO, blue) cell lines based on mRNA expression determined by RNA-sequencing.
**TRANSPARENT METHODS**

Cloning of CDK6 sgRNAs into px458 plasmid

pSpCas9(BB)-2A-GFP (px458) (plasmid number 48138; Addgene) was a gift from Krzysztof Chylinski. Cloning into px458 was performed as reported previously with slight modifications (Ran et al., 2013). The px458 plasmid was digested with BbsI (Thermofisher Scientific, Rockford, IL, USA) for 30 min at 37°C and gel purified using a Gel Extraction Kit (Qiagen, Hilden, Germany). The forward and reverse oligonucleotides (oligos) of each sgRNA were diluted at 100 μM in H2O. The CDK6 guide oligos contain overhangs for ligation into the BbsI sites in pSpCas9(BB) forward CDK6 oligo: CACCGCCCGGACTTTGAAGAACC, reverse CDK6 oligo: AAACCCGTCTCTGAGTCGGG. To phosphorylate and anneal the oligos, 2 μg of each oligo were mixed with T4 ligation buffer and T4 PNK (New England Biolabs, NEB, Ipswich, MA, USA) to a final volume of 20 μl and incubated for 1h at 37°C (phosphorylation), followed by 5 min at 95°C and then ramping down the temperature to 20°C at -1°C/min (annealing). Annealed and phosphorylated oligos were diluted 1:10 in H2O. Ligation reactions for each sgRNA were performed by mixing 10 ng of the digested and purified px458 plasmid with 90 ng of the diluted phosphorylated and annealed oligos, T4 ligation buffer, and T4 ligase (NEB) in a final volume of 10 μl. Ligation was carried out for 16 h at 16°C. Chemically competent DH5 bacteria were transformed and plated on LB plates containing 50 mg/ml ampicillin. Plates were incubated overnight at 37°C. Colonies were checked for correct insertion of the sgRNA by PCR colony screening, followed by sequencing (U6 primer: GAGGGCCTATTTCCCATGCATTAC).

Gene editing in BCR-ABL + cell lines

BCR-ABL+ cell lines were maintained in RPMI medium supplemented with 10% FCS, 50 μmol/L 2-mercaptoethanol, 100 U/mL penicillin, and 100 μg/mL streptomycin (PAA, Linz, Austria). Cells were electroporated (one single pulse with a voltage of 1600 V and a pulse width of 20 ms; NeonTM Transfection System, Invitrogen, Carlsbad, CA, USA) with 1μg control plasmid pSpCas9(BB)-2A-GFP(px458) or the plasmid expressing sgRNAs against CDK6 pSpCas9(BB)-2A-GFP(px458)-hCDK6del. For each transfection, 2x10^5 cells were resuspended in Buffer R (NeonTM) in a reaction volume of 10μl. Transfected cells were cultured in RPMI (supplemented with 10% FCS and 50 μmol/L 2-mercaptoethanol) for 24h. GFP+ cells were sorted using a FACSAriaTM II cell sorter (BD Biosciences, San Jose, CA, USA) and plated as single cell suspension into 96-well plates. For the re-expression of HA-CDK6, pMSCV-ires-GFP plasmids were used. The outgrowth of clones was monitored by microscopic inspection. Cells are routinely tested for mycoplasma contamination.

RNA extraction and Real-Time PCR

Total RNA was extracted from cell lines using the RNeasy Mini Kit (Qiagen) as recommended by the manufacturer. Reverse transcription was performed using the iSCRIPT cDNA synthesis kit (Bio-Rad). All qPCRs were performed in duplicate with the SsoFastTM EvaGreen®Supermix (Bio-Rad) according to the instructions of the manufacturer. Primer sequences for Dnmt3b were as follows: fwd, 5'-GTGGTGCACTGAGCTCGAAAGGATCTTCG-3' and rev, 5'-CACCGCCCGGACTTTGAAGAACC-3' (Kweon et al., 2017). Levels of mRNAs were normalized to Rplp0 mRNA.

Western Blotting

Protein lysates were prepared using RIPA lysis buffer supplemented with cComplete™ Protease Inhibitor tablet (Roche Diagnostics, Indianapolis, IN, USA), separated by SDS/PAGE and transferred onto nitrocellulose blotting membrane (Amersharm™ Protran™ 0.45μm NC, GE Healthcare, Life Sciences). Membranes were blocked in 5% BSA followed by incubation with primary antibodies: anti-CDK6 (1:1000, H-96, Santa Cruz, Dallas, TX, USA), anti-HA (1:1000, ab9110, Abcam, Cambridge, UK), anti-Dnmt3b (1:1000, ab122932, Abcam) and anti-HSC70 (1:1000, B-6, Santa Cruz). Appropriate secondary HRP antibodies (1:10000, Cell Signaling Technology, Danvers, MA, USA) were used and membranes were visualized using Clarity™ ECL Western blotting substrate (Biorad, Hercules, CA, USA).

Reduced Representation Bisulfite Sequencing (RRBS)

RRBS was performed as reported previously (Heller et al., 2016). In brief, genomic DNA was MspI (NEB) digested, end-repaired and A-tailed using Klenow Polymerase (NEB) followed by adapter ligation using Quick Ligase (NEB) and AMPure XP size selection (Beckman Coulter, Fullerton, CA, USA). RRBS libraries were sodium bisulfite treated using the EZ-DNA Methylation-Direct kit (Zymo Research Corp, Orange, CA, USA) and quantified by qPCR. Enrichment PCR was performed using the PfuTurboCx Hotstart Kit (Agilent Technologies, Santa Clara, CA, USA) followed by AMPure XP clean up. Quality of final RRBS libraries was determined by Experion analysis (Biorad). Sequencing was performed on a HiSeq2000 sequencer (Illumina Inc, San Diego, CA, USA). RRBS reads were processed using the --
platform, utilizing the Sequest described via RP pH 8 and all used for offline fractionation for the analysis of the phosphoproteome. Offline fractionation TFA, phosphopeptides were eluted with 0.01% µg TMT with a fresh solution of Fe(III)Cl trifluoroacetic to achieve a final peptide concentration of ~1µg/µl. Eluate was then used for a reaction as previously described performed for 30 min at 37°C. The optimum number of amplification cycles was estimated by qPCR reaction as previously described (Rendeiro et al., 2016). Following library amplification, fragments larger than 1 200 bp were excluded by SPRI size selection. DNA concentration was measured using a Qubit fluorometer (Invitrogen). Libraries were amplified using custom Nextera primers (Buenrostro et al., 2013) and sequenced by the Biomedical Sequencing Facility at CeMM using the Illumina HiSeq3000/4000 platform. ATAC-seq data are deposited in the Gene Expression Omnibus ( GEO) database (Accession ID: GSE113752).

ChIP-seq
CDK6 chromatin immunoprecipitation (ChIP) was performed using an antibody against HA (ab9110, Abcam) as described previously (Bellutti et al., 2018, Scheicher et al., 2015). Cells were crosslinked with DSG (20 minutes, RT) and 1% formaldehyde (10 minutes, RT) and termination of the fixation procedure was performed using glycine. For immunoprecipitation (IP), 70 µl Dynabeads Protein G magnetic beads (Invitrogen) were used, IP DNA was subjected to sequencing library preparation and sequencing was performed using the Illumina HiSeq3000/4000 platform. Raw sequencing reads were quality controlled using FASTQC followed by quality filtering, trimming of reads and adapter removal using trimmomatic (v0.36). Mapping against the mouse reference genome (Gencode M13) was done using bwa mem (v0.7.15) and blacklisted regions were removed using bedtools subtract (v2.26.0). Multimappers and reads with bad mapping quality were removed using samtools (v1.3.1). Peak calling was performed using MACS2 (v2.1.0) using default parameters and motif identification was performed using Homer (v4.9.0) findMotifsGenome.pl with the default -size 200. ChIP-seq data are deposited in the Gene Expression Omnibus ( GEO) database (Accession ID: GSE156966).

ATAC-seq
10^6 cells were washed once in 50 µl PBS and resuspended in transposase reaction mix (12.5 µl 2 × TD buffer, 2 µl transposase (Illumina), 10.5 µl nuclease-free water and 0.01% NP-40). Tagmentation was performed for 30 min at 37°C. Following library amplification, fragments larger than 1 200 bp were excluded by SPRI size selection. DNA concentration was measured using a Qubit fluorometer (Invitrogen). Libraries were amplified using custom Nextera primers (Buenrostro et al., 2013) and sequenced by the Biomedical Sequencing Facility at CeMM using the Illumina HiSeq3000/4000 platform. ATAC-seq data are deposited in the Gene Expression Omnibus ( GEO) database (Accession ID: GSE156966).

Chromatome analysis
Chromatome analyses were performed as described previously (Dutta et al., 2012, Bellutti et al., 2018). Briefly, the purified chromatin pellet was subjected to Benzonase digestion and solubilized in SDS lysis buffer. Filter Aided Sample Prep (FASP) was performed according to the procedure described previously (Wisniewski et al., 2009). Peptides were desalted using C18 solid phase extraction spin columns (The Nest Group, Southborough, MA) labeled with TMT 6plex™ reagents (Pierce, Rockford, IL) and pooled. Organic solvent was removed in vacuum concentrator and labelled peptides were loaded onto a solid phase extraction column. Peptides were eluted with 300µL 80% acetonitrile containing 0.1% trifluoroacetic acid to achieve a final peptide concentration of ~1µg/µl. Eluate was then used for phosphopeptide enrichment applying a modified method of immobilized metal affinity chromatography (IMAC) (Ficarro et al., 2005). Briefly, two times 100 µl of Ni-NTA superflo slurry (Qiagen) were washed with LCMS-grade water and Ni³⁺ stripped off the beads by incubation with 100 mM of EDTA, pH 8 solution for 1 h at room temperature. Stripped NTA resin was recharged with Fe⁶⁺-ions by incubation with a fresh solution of Fe(III)Cl₃ and 75 µL of charged resin used for the enrichment of a total of ~300 µg TMT-labelled peptide. The unbound fraction was transferred to a fresh glass vial and used for offline fractionation for the analysis of the whole chromatome proteome. After washing the slurry with 0.1% TFA, phosphopeptides were eluted with a freshly prepared ammonia solution containing 3mM EDTA, pH 8 and all used for offline fractionation for the analysis of the phosphoproteome. Offline fractionation via RP-HPLC at pH 10 and 2D-RP/RP Liquid Chromatography Mass Spectrometry were performed as described (Bellutti et al., 2018). Raw data files were processed using the Proteome Discoverer 2.2.0. platform, utilizing the Sequest HT database search engine and Percolator validation software node.
(V3.04) to remove false positives with a false discovery rate (FDR) of 1% on peptide and protein level under strict conditions. Searches were performed with full tryptic digestion against the mouse SwissProt database v2017.12 (25 293 sequences and appended known contaminants) with up to two miscleavage sites (Bellutti et al., 2018). For statistical analysis and p-value calculation, the integrated ANOVA hypothesis test was used. TMT ratios with p-values below 0.01 were considered statistically significant.

Publicly available datasets and co-expression analysis
The following human ALL Affymetrix HG-U133_plus_2.0 gene expression microarray datasets were obtained from ArrayExpress database: E-MTAB-5035, E-GEOD-13351, E-GEOD-49032 and E-GEOD-13576. Spearman correlation coefficients of log2 expression values were calculated using the “cor” function of R and data visualization was done using the “corrplot” package. ETS1 ChIP-seq data were obtained from GEO database: GSM803442, GSM803468, GSE83758. Infinium Human Methylation 450K BeadChip data of AML patients were obtained from the Cancer Genome Atlas database (LAML dataset).
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