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**PHF6-altered T-ALL harbored epigenetic repressive switch at bivalent promoters and respond to 5-azacitidine and venetoclax**

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**Running title :**

PHF6<sup>ALT</sup> T-ALL respond to 5-azacitidine and venetoclax

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**Conflicts of interest :**

The authors have declared that no conflict of interest exists.

## **Abstract**

**Purpose:** To assess the impact of PHF6 alterations on clinical outcome and therapeutical actionability in T cells acute lymphoblastic leukemia (T-ALL).

**Experimental Design:** We described PHF6 alterations in an adult cohort of T-ALL from the French trial GRAALL 2003/2005 and retrospectively analyzed clinical outcomes between PHF6-altered (PHF6<sup>ALT</sup>) and wild-type patients. We also used EPIC and ChIP-seq data of patient samples to analyze the epigenetic landscape of PHF6<sup>ALT</sup> T-ALLs. We consecutively evaluated 5-azacitidine efficacy, alone or combine with venetoclax, in PHF6<sup>ALT</sup> T-ALL.

**Results:** We show that PHF6 alterations account for 47% of cases in our cohort and demonstrate that PHF6<sup>ALT</sup> T-ALL presented significantly better clinical outcomes. Integrative analysis of DNA methylation and histone marks shows that PHF6<sup>ALT</sup> are characterized by DNA hypermethylation and H3K27me3 loss at promoters physiologically bivalent in thymocytes. Using patient-derived xenografts (PDX), we show that PHF6<sup>ALT</sup> T-ALL respond to the 5-azacytidine alone. Finally, synergism with the BCL2-inhibitor venetoclax was demonstrated in refractory/relapsing PHF6<sup>ALT</sup> T-ALL using fresh samples. Importantly, we report three cases of refractory/relapsed (R/R) PHF6<sup>ALT</sup> patients who were successfully treated with this combination.

**Conclusions:** Overall, our study supports the use of PHF6 alterations as a biomarker of sensitivity to 5-azacytidine and venetoclax combination in R/R T-ALL.

## **Translational Relevance**

Predicting and treating R/R T-ALL remains a major challenge and require improved patient prognosis stratification and biomarkers for targeted therapies. PHF6 is frequently mutated in adult T-ALL, but the impact of its alterations on clinical outcome and their actionability as a therapeutic target remain elusive. We describe PHF6 alterations in 216 T-ALL and show that PHF6 loss of function is associated with better outcome, independently of other known prognostic parameters. Using multiomics approaches, we demonstrate that PHF6 alterations in T-ALL is associated with DNA hypermethylation and H3K27me3 loss at promoters physiologically bivalent in thymocytes. Using fresh samples of R/R T-ALL, we revealed increased in vivo sensitivity of PHF6<sup>ALT</sup> T-ALL to the demethylating agent 5-azacitidine alone or associated with the BCL2-inhibitor venetoclax. We report three cases of R/R PHF6<sup>ALT</sup> T-ALL patients successfully treated with this combination. Overall, our study supports the use of PHF6 alterations as a biomarker of eligibility to 5-azacytidine and venetoclax in R/R T-ALL.

## Introduction

T-cell acute lymphoblastic leukemia (T-ALL) is a heterogeneous hematological disease resulting from malignant proliferation of immature thymic cell precursors blocked at varying stages of differentiation, which partially recapitulates normal lymphoid T-cell ontogeny (1). T-ALL mostly occurs in children and young adults (10%-15% and 20%-25% of ALL cases, respectively) (2). The use of intensive chemotherapy regimens has considerably improved the outcome of adult T-ALL. Yet, refractory and relapsed disease conveys a dismal prognosis with a 6-to-9 months median survival (2–5). Therefore, improving patient prognosis stratification and finding biomarkers for innovative targeted therapies is of major importance. Epigenetic alterations are frequent events in T-ALL and can offer unique therapeutic vulnerabilities to epigenetic modifiers such as the hypomethylating agent 5-azacytidine. However, the eligibility of T-ALL patients to this therapy is still debated (6,7).

Plant Homeodomain Finger protein 6 gene (*PHF6*) was originally identified as the causative gene of Börjesson-Forsmann-Lehman syndrome (BFLS), a rare X-linked neurodevelopmental disorder (8). *PHF6* is frequently altered by somatic LoF mutations and deletions in T-ALL (~40% adult and ~15% pediatric cases) and T/M mixed-phenotype acute leukemia (MPAL), and less frequently in myeloid malignancies or clonal hematopoiesis (9–14). The PHF6 protein localizes within the nucleolus where it support a role in the control of transcription and ribosome biogenesis (15,16). In addition, PHF6 has been demonstrated to control chromatin accessibility through interaction with chromatin remodelers such as NuRD and SWI/SNF complexes (15,17–19).

*PHF6* alterations are associated with NOTCH1 and IL7R/JAK/STAT pathway genes gain-of-function mutations, PRC2 complex genes LoF mutations, *WT1* and *PTPN2*

deletions and TLX1 or TLX3 ectopic expression (20–27). Analysis of clonal evolution and mutation dynamics using paired diagnostic and relapse samples identified *PHF6* LoF as an early events in T leukemogenesis (28). Mechanistically, increased self-renewal in the hematopoietic stem cell compartment following *PHF6* LoF have been reported as a major feature contributing to leukemia (28–31). Even though *PHF6* alterations are often found in adult T-ALL, their impact on prognosis and their therapeutic actionability remain poorly explored (31–33).

In this study, we analyzed *PHF6* alterations in a large cohort of adult T-ALL from the Group for Research on Adult Acute Lymphoblastic Leukemia (GRAALL) 03/05 French trials. Our data demonstrate that *PHF6* alterations in T-ALL lead to DNA hypermethylation and is associated with response to demethylating drugs alone or in combination with the BCL2 inhibitor venetoclax.

## **Material and Methods**

### **Patients enrolled in the GRAALL 2003-2005 study**

Patients were enrolled in the GRAALL-2003-2005 trials (registered on <http://clinicaltrials.gov> as follows: GRAALL-2003, #NCT00222027; GRAALL-2005, #NCT00327678). 216 patients out of 337 were included in this study based on DNA availability for molecular analyses. No differences in clinical outcomes were observed between the included patients and the entire cohort (data not shown). Diagnostic samples were collected after informed consent according to the Declaration of Helsinki. Immunophenotyping was performed as previously described (1). All samples used contained at least 80% blasts. Patient minimal residual disease (MRD) was assessed as previously described (59).

### **Patients enrolled in the ALL-TARGET study**

The three 5-AZA-Venetoclax treated-patients reported in this manuscript have been registered within the ALL-TARGET, Registry of Relapsed/Refractory T-cell Acute Lymphoblastic Leukemia, trial. This study has been registered on ClinicalTrials.gov. under NCT05832125 references.

### **Next-generation sequencing, copy number, and molecular analyses**

Genomic analysis was performed by pan-exon targeted next-generation sequencing of DNA extracted from diagnosis samples. The next-generation sequencing panel included 103 genes relevant in hematological malignancies (Nextera XT gene panel; Illumina, San Diego, CA). Libraries were prepared according to Illumina instructions and sequenced on a MiSeq instrument (Illumina; 500x with mean coverage 95%). Sequence reads were aligned to the reference genome (GRCh38) using our in-house software Polyweb (Institut Imagine, Paris). Variant filtering and calling were performed using validated criteria (coverage <30x, <10 alternative reads or variant allelic fraction < 7%, polymorphisms described in dbSNP, 1000Genomes, EVS, Gnomad, and EXAC with a calculated mean population frequency >0.1%); annotations were done using somatic database COSMIC and ProteinPaint (St Jude Children's Research Hospital—Pediatric Cancer data portal). Some mutations were confirmed by Sanger sequencing. Copy number variants were assessed either by multiplex ligation-dependent probe amplification technique (SALSA-multiplex ligation-dependent probe amplification P383 T-ALL probe mix (MRC-Holland, Amsterdam, Netherlands) kit containing 53 probes for 13 different chromosomal regions of diagnostic or prognostic importance in T-ALL. PHF6 deletions were confirmed by high-resolution array comparative genomic hybridization as previously reported (35).

### **EPIC array analysis**

Intensity Data (IDAT) files from Illumina Infinium Methylation EPIC BeadChip (EPIC arrays) were processed and normalized with RnBeads Bioconductor package. Probes with high SNP probability and bad detection P-values were filtered out. EPIC arrays were also used for copy number analysis using the conumee Bioconductor package, and further processed with GISTIC2 to identify recurrent copy number aberrations. Probes located in regions with altered copy numbers were excluded from downstream analysis. Differential methylation analysis was performed using Limma R package (60).

### **ChIP-seq analysis**

BAM files for thymocytes ChIP-seq datasets were obtained from the BLUEPRINT consortium. Peak calling was performed as previously described (38). Regulatory features were found by overlapping histone marks peaks with the GenomicRanges R package. Enrichment analysis of DMPs within regulatory regions was performed with the LOLA Bioconductor package. As background probes, we used all the annotated probes of the EPIC analysis.

### **RNA sequencing and data analysis**

For T-ALL, 89 primary samples with eligible RNA were analyzed by poly(A)-enriched RNA-Seq. Fragments were sequenced in stranded paired-end mode (2×50 bp) using the Agilent platform. Variance-stabilized log<sub>2</sub> expression values were computed using DESeq2 R package. Z-scores of log<sub>2</sub> expression values were computed to perform hierarchical clustering analysis using Euclidean distance and complete clustering method arguments with ComplexHeatmap R package. Uniform Manifold Approximation and Projection for dimension reduction was performed using the M3C R package.



### **Patient-derived xenografts**

Patient-derived xenografts (PDX) were generated from primary T-ALL samples as previously reported.<sup>38</sup>  $10^6$  viable leukemic cells were xenografted by intravenous retro-orbital injection in 6-weeks old NSG mice. Mice were monitored weekly by flow cytometry for leukemic load (FSChi, hCD7+, hCD45+ cells) in peripheral blood. Animals were clinically and biologically monitored until endpoint was reached or terminally ill, according to local ethical rules and home office license (30078-2021021814199445). Bone marrows from tibiae, hip, femora and vertebrae were collected for subsequent ex vivo experiments. All samples used contained  $\geq 90\%$  blasts.

### **PDX 5-azacytidine treatment**

Mice were treated in a “curative-like” setting, starting when peripheral blood blast counts exceeded 1%. They received 5 mg/kg/day 5-Azacytidine intra-peritoneally (IP) for five days twice with two days break. Untreated mice received the same volume of NaCl IP. Mice were sacrificed when the blast fraction was more than 80% or if mice presented clinical signs of disease (loss of weight  $>10\%$ , neurological symptoms, tumor development). The national ethics committee approved the mouse study: PROJET APAFIS # 8853 N° 2017020814103710.

### **Cytotoxic assay of fresh patient samples**

Fresh blast-infiltrated samples were counted for cellularity, assayed by flow cytometry to quantify blast-infiltration. Samples had at least  $20 \cdot 10^6$  cells with more than 25% blasts and 50% of viability before incubation with drugs. Cell suspension was incubated with 1.25 to 5  $\mu\text{g}/\mu\text{L}$  of 5-azacytidine combined, or not with 125  $\mu\text{M}$  of venetoclax. Cells were then cultured in a complete medium supplemented with cytokines, at 37°C. After 3 days of incubation, cell viability was assessed by flow cytometry (Annexin-V/Propidium iodide).

## Statistical analyses

Statistical analyses were performed using GraphPad Prism 8 software and R (version 4.2.0). The following were used to indicate significant differences: • :  $P > 0.05$ ; \* :  $P < 0.05$ ; \*\* :  $P < 0.01$ ; \*\*\* :  $P < 0.001$ ; \*\*\*\* :  $P < 0.0001$ .

## Data availability

The data generated in this study are not publicly available due to information that could compromise patient privacy or consent but are available upon reasonable request from the corresponding author, [vahid.asnafi@aphp.fr](mailto:vahid.asnafi@aphp.fr).

## Results

### ***PHF6* alterations are frequent in adult T-ALL**

To assess the prevalence of *PHF6* alterations in T-ALL, we analyzed data from targeted whole-exome sequencing and copy number analysis previously obtained for 216 adult patients from the GRAALL-03/05 protocols (34). 47% patients had at least one *PHF6* LoF alteration in our cohort (102 ALT patients versus 114 WT) (Figure 1A). The most frequent alterations were nonsense mutations followed by frameshift indels and missense mutations (Figure 1B-C). Truncating mutations were frequently located within the 5' half of the protein whereas missense mutations were almost exclusively found at the ePHD2 and 3' PHD zinc finger domain.

We then compared the oncogenetic landscape between *PHF6*<sup>ALT</sup> and *PHF6*<sup>WT</sup> T-ALL. *PHF6* LoF was significantly associated with increased mutational rates of *WT1* and *RUNX1*, and decreased mutational rate of *ETV6*, three transcription factors known to be frequently altered in immature T-ALL. As previously reported, *PHF6* alterations frequently co-occurred with activating NOTCH1 and IL7R pathway (*IL7R*, *JAK1*, *JAK3*, *STAT5B*, *DNM2*, *PTPRC*, *PTPN2*) mutations. The RAS signaling regulator NF1 was

also frequently co-inactivated. Among epigenetic factors, *PHF6* alterations were associated with increased mutational rates of the PRC2 complex gene *SUZ12*, *ASXL1* and *CTCF*. Conversely, *PHF6* alterations segregated from mutations in *DNMT3A* and *IDH2*, both implicated in DNA methylation process (Figure 1D-E and Supplementary Figure S1A). *PHF6* alterations were less frequent in mature TCRAB expressing T-ALL (25% vs 75% with  $p = 0.01$ ) (Figure 1F). Consistent with seminal studies, *PHF6* LoF anomalies were significantly associated with ectopic expression of both TLX1 and TLX3 with marked enrichment in the TLX1 subtype (35% vs 12% with  $p < 1.10^{-4}$  and 19% vs 9% with  $p = 0.03$  respectively) (Figure 1G). In line with their absence in TCRAB expressing T-ALL, *PHF6* alterations were virtually absent from the TAL1 deregulated subtype (1% in *PHF6*<sup>ALT</sup> T-ALL vs 26% in *PHF6*<sup>WT</sup> cases,  $p < 1.10^{-4}$ ).

Finally, we addressed the question of the conservation of *PHF6* alterations from diagnosis to relapse. In a series of 5 protocolar paired diagnosis/relapsed *PHF6*<sup>ALT</sup> patients, all retained *PHF6* alteration at relapse (Supplementary Figure S1B). We decided to confirm these results on a larger non protocolar cohort of 37 other patients bearing *PHF6* pathogenic variant at diagnosis (Supplementary Table 1). Among them, 35 (95%) retained *PHF6* alteration at relapse. These results suggest that *PHF6* alterations are strongly conserved during the clonal evolution of the disease.

### ***PHF6* alterations are associated with better clinical outcomes in the GRAALL-03/05 protocols**

A clinical and biological comparison of cases with and without *PHF6* alterations is shown in Table 1. *PHF6*<sup>ALT</sup> patients were significantly older than the rest of the cohort (median age of 34.17 versus 28.22 years with  $p = 0.04$ ).

We also analyzed treatment response in *PHF6*<sup>ALT</sup> and *PHF6*<sup>WT</sup> cases in our cohort. There was no significant difference between the two groups regarding steroid response and early bone marrow blast clearance. Nevertheless, and despite older age, *PHF6*<sup>ALT</sup> patients had significantly lower rates of death during induction (1% versus 7%,  $p = 0.04$ ), and higher attainment of complete remission (96% versus 89%,  $p = 0.05$ ). Mutated patients had better outcomes at five years estimation in terms of overall survival, event-free survival, and cumulative incidence of relapse.

By univariate analysis, *PHF6*<sup>ALT</sup> patients had significantly increased overall survival (HR 0.51,  $p = 0.004$ ) (Table 1 and Figure 2). This benefit remained significant by multivariate analysis and was independent from the High risk genetic classifier (HR of 0.58 with  $p = 0.03$ ) (35). Overall, our results show that *PHF6* LoF mutations are associated with better treatment response and survival and should be considered when assessing patient prognosis.

### ***PHF6*<sup>ALT</sup> T-ALL display DNA hypermethylation at promoters of physiological PRC2 target genes**

*PHF6* being a known epigenetic modulator, we sought to explore the epigenetic landscape of *PHF6*<sup>ALT</sup> T-ALL. We thus performed genome-wide methylation analysis, taking advantage of data formerly generated by our team, in a series of 143 primary adult T-ALL samples (7). Genome-wide DNA methylation was significantly higher in

*PHF6*<sup>ALT</sup> when compared to WT patients (Figure 3A). These results are coherent with our previous paper on DNA methylation in T-ALL in which *PHF6* alterations seemed to be associated with hypermethylated cluster C3, C4 and C5 (7). Indeed *PHF6* alterations were significantly enriched in clusters C3 and C4 (15/22,  $p = 0.01$  and 30/41,  $p < 0.001$  respectively) and almost absent from C1 and C2 hypomethylated clusters (0/14,  $p < 0.001$  and 1/34,  $p < 0.001$  respectively)(Supplementary Figure S2A). There was no significant enrichment for *PHF6* alterations in cluster C5 (16/32,  $p = 0.25$ ). This cluster was reported to have dismal outcomes in our previous paper. We thus checked if *PHF6* status could refine the overall survival prognosis in this cluster. Interestingly yet not significantly, *PHF6*<sup>ALT</sup> patients from the C5 cluster had a tendency towards better overall survival than their WT counterpart (Supplementary Figure S2B).

Differential methylation analysis identified a total of 42,164 differentially methylated probes (DMP) (39,286 hypermethylated and only 2,878 hypomethylated DMP) in *PHF6*<sup>ALT</sup> T-ALL when compared to *PHF6*<sup>WT</sup> (Figure 3B). Hypermethylated DMPs (hyDMPs) were strongly enriched in CpG islands (CGIs), regardless of their genomic location. Hypomethylated DMPs were enriched in CpG open seas and shores and preferentially located at distal regions of promoter and gene body (Figure 3C and 4D). Of note, hyDMPs were strictly hypomethylated in normal thymocytes (Figure 3E). A total of 3,066 protein-coding gene-associated promoters were hypermethylated and 431 were hypomethylated in *PHF6*<sup>ALT</sup> T-ALL (Figure 3F). Hypermethylated promoters were associated with H3K27me3 and H3K4me3 histone marks in CD34+ and CD3+ cells (Figure 3G). ChIP-seq analysis of these promoters at different stages of normal thymocytes revealed simultaneous H3K4me3 and H3K27me3 marks, representing bivalent chromatin domains, a hallmark of developmentally regulated genes (Figure 3H) (36,37). Consistently, these promoters were enriched in target genes of SUZ12

and EZH2, two subunits of the PRC2 complex, responsible for H3K27me3 histone mark deposition and a major regulator of bivalent chromatin.

*PHF6* and *SUZ12* alterations significantly co-occur in our cohorts, therefore, to prevent potential confounding factors, patients were separated into *PHF6* altered only, PRC2 altered only (with alterations of *SUZ12*, *EED* or *EZH2*), both altered, or none altered. T-ALL bearing alterations in either *PHF6*, PRC2 or in both, had significantly higher overall DNA methylation (Figure 3I). These T-ALL shared 30,839 hyDMPs when compared to T-ALL with no alterations and constitute a coherent cluster on UMAP projection of methylome profile (Figure 3J and K). These hyDMPs were also enriched at genes associated with H3K27me3 and H3K4me3 histone marks in CD34+ and CD3+ cells and affiliated with gene ontology terms related to developmental processes, transcription regulation, cell proliferation and central nervous system differentiation (Figure 3L). Altogether, our data suggest that *PHF6*<sup>ALT</sup> T-ALL display DNA hypermethylation at promoters of physiological PRC2 target genes.

### ***PHF6*<sup>ALT</sup> T-ALL display altered epigenetics landscape of bivalent promoters with loss of H3K27me3 and DNA hypermethylation**

Given the enrichment of hyDMPs in both H3K27me3 and H3K4me1/3, we hypothesized that *PHF6*<sup>ALT</sup> T-ALL display aberrant DNA hypermethylation and profound alteration of bivalent promoters.

We therefore performed enrichment analysis for regulatory elements on hyDMPs using ChIP-seq data of H3K4me3 and H3K27me3 histone marks. Two types of promoters were defined and studied (active and bivalent) (38). To assess the possible implication of major oncotypes we also studied the distribution of TLX1, TAL1 and PHF6 binding sites by using publicly available ChIP-seq data (39). hyDMPs and PHF6-bound regions

were strikingly enriched in bivalent promoters. Of note, among known genes targeted by PHF6 (118 genes), 11 (~10%) are also found in repressed bivalent genes (Supplementary Table 2). Genomic regions bound by TAL1 or TLX1 did not overlap with bivalent promoters but exclusively with active one (Figure 4A). Unsupervised hierarchical clustering of DNA methylation level efficiently segregated samples according to their oncotype when considering active promoters, while bivalent ones separated *PHF6*<sup>ALT</sup> T-ALL from *PHF6*<sup>WT</sup> T-ALL and normal thymocytes, regardless of the oncotype (Figure 4B). Strikingly, a subset of *PHF6*<sup>WT</sup> T-ALL, strongly enriched in TAL1 oncotype showed very similar methylation profile of bivalent promoters to normal thymocytes, with very low levels of DNA methylation. UMAP projection according to bivalent promoters methylation levels displayed similar results (Figure 4C). Hypermethylation of bivalent regions has already been described and usually follows the loss of bivalent histone marks (40–42). To look for epigenetic changes in histone mark deposition following *PHF6* LoF, we analyzed H3K27me3 and H3K4me3 ChIP-seq peaks from 8 T-ALL (4 *PHF6*<sup>WT</sup>/*PRC2*<sup>WT</sup>, 2 *PHF6*<sup>ALT</sup>, *PRC2*<sup>ALT</sup>, *PHF6*<sup>ALT</sup>/*PRC2*<sup>ALT</sup>) at thymic bivalent promoters. *PHF6*<sup>ALT</sup> T-ALL had a profound reduction of H3K27me3 peaks at thymic bivalent promoters when compared with WT T-ALL (Figure 4D). Among the 2278 thymic bivalent genes, 1545 were associated to hyDMPs (Figure 4E). Surprisingly, *PRC2*<sup>ALT</sup> T-ALL had no reduction of H3K27me3 peaks while *PHF6*<sup>ALT</sup>/*PRC2*<sup>ALT</sup> had profiles similar to *PHF6*<sup>ALT</sup>. The analysis of H3K4me3 marks gave comparable results, although less pronounced. These results suggest a role for PHF6 in the maintenance of H3K27me3 histone mark and of low levels of DNA methylation at bivalent promoters.

We then sought to assess hyDMRs-associated bivalent gene expression in T-ALL and look for differential expression between *PHF6*<sup>ALT</sup> and *PHF6*<sup>WT</sup> samples. We therefore

performed differential expression analysis using bulk RNA-seq on 89/216 primary clinical samples (49 WT and 40 ALT). Out of the 1548 bivalent genes associated with hyDMRs, only 65 were differentially expressed between the two conditions (27 upregulated and 38 downregulated in *PHF6*<sup>ALT</sup> T-ALL), suggesting epigenetic alteration of bivalent promoter has minimal impact on bivalent genes expression at the stage of patent leukemia (Figure 4F).

### ***PHF6*<sup>ALT</sup> T-ALL are sensitive to hypomethylating agents alone or in combination with BCL2 inhibitor venetoclax**

*PHF6*<sup>ALT</sup> T-ALL have significantly higher level of global DNA methylation. We thus reasoned that treatment with hypomethylating agents could be of potential interest in the treatment of these T-ALL. To first confirm this rational, we took advantage of results previously generated with patient derived xenografts (PDX) from 4 primary T-ALL samples (2 *PHF6* WT (UPNT-M525 / UPNT-M894) and 3 ALT (UPNT-M149 / UPNT-670)) (7). Treatment was initiated when peripheral blood (PB) blast counts exceeded 1%. Mice were given 5-azacytidine in vivo or left untreated. For PDX derived from *PHF6*<sup>ALT</sup> patients, curative 5-azacytidine treatment significantly increased survival (Figure 5A). There were no differences in terms of survival between treated and untreated mice derived from *PHF6*<sup>WT</sup> patients-derived PDX. These results suggest that *PHF6*<sup>ALT</sup> T-ALL indeed display increased vulnerability to demethylating agents.

R/R disease being a major problem in T-ALL, due to particularly poor prognosis, we examined whether 5-azacytidine could be used to treat patients at relapse. We therefore performed ex vivo cytotoxic assays using 5-azacytidine on fresh blasts infiltrated samples from 21 R/R T-ALL patients collected within the French ALL-Target protocol (9 *PHF6*<sup>ALT</sup>, 5 *PRC2*<sup>ALT</sup>, 7 WT)(Supplementary Table 3). *PHF6*<sup>ALT</sup> samples had significantly better response to 5-azacytidine than WT (Figure 5B). Coherently with



their comparable DNA methylation profile, similar results could be observed for PRC2<sup>ALT</sup> T-ALL.

Anti-apoptotic dependency of T-ALL vary according to their phenotype with immature and early cortical T-ALL survival relying essentially on BCL2.<sup>56</sup> We thus wondered whether *PHF6* alterations could be associated with better response to 5-azacytidine and venetoclax combination, a reference treatment combination for AML uneligible for intensive chemotherapy and an emerging one for T-ALL (43–45). Of note, within GRAALL03/05 cohort, *PHF6*<sup>ALT</sup> T-ALL indeed displayed higher BCL2 transcript expression when compared to WT, further supporting this rationale (Figure 5C). We therefore tested this combination ex vivo, on fresh blasts from the same patients depicted in fig. 5B. There was no significant difference between *PHF6*<sup>ALT</sup> and WT samples when exposed to 125 nM of venetoclax. Nevertheless, *PHF6*<sup>ALT</sup> blasts were significantly more sensitive to the combination of both drugs when compared to WT (Figure 5D). Again, PRC2<sup>ALT</sup> T-ALL displayed similar response profile to this combination.

Given these results, we investigated the efficacy of this combination at relapse in a small cohort of patients with relapsed or refractory disease. Three R/R *PHF6*<sup>ALT</sup> patients (from those tested in vitro in Figure 5B and D, details are provided in Supplementary Table 3) were treated with 5-azacytidine and venetoclax combination at relapse (Figure 5E). Patient 1 (T-ALL1) presented with T lymphoblastic lymphoma at diagnosis and relapsed as a T-ALL during the first year of treatment according to the standard of care. He received a total of three lines of treatment before benefiting from 5-azacytidine and venetoclax combination. Complete bone marrow blast clearance and metabolic response, assessed by positron emission tomography (PET), was achieved after the first course of treatment. Patient 2 (T-ALL2) was treated with 5-

azacytidine and venetoclax upon mediastinal relapse 15 months after HSCT. Complete morphological and metabolic response (Deauville 4), was obtained after two courses of treatment. Patient 3 (T-ALL3) was treated with the same combination at relapse with cutaneous and CNS involvement, four years after HSCT. Complete medullary and extramedullary response was achieved after two courses and four intrathecal chemotherapies, allowing second HSCT. Apart from expected adverse events, treatment was well tolerated. All three patients are alive at the time of writing this manuscript. Taken together, these results suggest that *PHF6* LoF is associated to increased sensitivity to 5-azacytidine alone or in combination with venetoclax.

## **Discussion**

This study is the first to assess *PHF6* alterations incidence and clinical impact in a large protocolar study. Consistent with previous literature, *PHF6* alterations are frequent in GRAALL 2003-2005 cohort (10,20–27). As reported in previous studies, *PHF6* alterations are strongly associated to TLX oncotypes, with particular emphasis for TLX1. *PHF6* LoF seems to be mutually exclusive with a mature TCRAB phenotype in T-ALL and with known *TAL1* expression deregulations.

We revealed that *PHF6*<sup>ALT</sup> T-ALL are characterized by high overall DNA methylation. Differentially hypermethylated probes were highly enriched in CGIs located at regions physiologically bearing bivalent H3K4me3 and/or H3K27me3 histone marks. Such chromatin is frequently associated with genes regulated by Polycomb complexes and H3K4me3 depositing proteins such as COMPASS and MLL2 (KMT2B/D) complexes (37,46,47). These genes are involved in developmental processes and thought to display highly variable and finely tuned expression. Moreover, they have been implicated in the regulation of lineage specificity and functional plasticity of differentiating CD4<sup>+</sup> T cells, revealing their importance during thymocytes maturation

(48). Many bivalent genes in human ES cells (hESC) are frequent targets for hypermethylation in human cancers and growing evidence suggests a role in carcinogenesis can be found in literature (40,46,49–51). Our results show that thymic bivalent promoters are maintained strictly hypomethylated during thymopoiesis. Even though hypermethylation of these promoters had little impact on bivalent genes expression in *PHF6<sup>ALT</sup>* T-ALL, loss of PHF6 and subsequent modification of the epigenetic landscape could participate to leukemogenesis by altering thymopoiesis. Supporting a role for PHF6 in T cell development, mouse models did find altered thymopoiesis with reduced DN2 and DN3 thymocytes, as well as lower counts of T cells in peripheral blood after conditional *Phf6* KO (28,30,52).

PRC2 is thought to protect CGIs from methylation, directly or through the recruitment of other epigenetic regulators like TET proteins (41,42,53). DNA hypermethylation of PRC2 target genes have already been described in T-ALL (42,54). Our results indicate that *PHF6* and/or PRC2-altered T-ALL display a switch from a labile and permissive silencing of bivalent genes, insured by H3K27me3, to a more static state following DNA hypermethylation (55). We propose that both *PHF6* and PRC2 complex are implicated in the maintenance of thymic bivalent regions and might protect them from DNA methylation in a similar manner. Mutual exclusion of *PHF6* and *DNMT3A* LoF suggests that this hypermethylation is crucial during T leukemogenesis. Notably, *PHF6* locates at bivalent promoters and has been shown to interact with *RBBP4*, a member of PRC2, *MLL2* and *NuRD* complexes, all implicated in bivalent promoters maintenance (47,56,57). Moreover, the co-occurrence of *PHF6* and PRC2 alterations support a collaborative interaction during leukemogenesis. Surprisingly, *PHF6<sup>ALT</sup>/PRC2<sup>WT</sup>* T-ALL only showed a drastic reduction in H3K27me3 mark at bivalent promoters while *PHF6<sup>WT</sup>/PRC2<sup>ALT</sup>* had almost none. These results could be explained by residual

activity of the PRC2 complex due to sufficient expression of the remaining wild type allele in *PHF6*<sup>WT</sup>/*PRC2*<sup>ALT</sup> T-ALL. Because of its location on X chromosome, *PHF6* is more prone to permanent LoF when mutated, even in the case of female patient, as a result of X inactivation. However, both *PHF6*<sup>ALT</sup>/*PRC2*<sup>WT</sup> and *PHF6*<sup>WT</sup>/*PRC2*<sup>ALT</sup> T-ALL display hypermethylation of bivalent promoters suggesting that loss of H3K27me3 is dispensable during this process. Overall, this study supports a model in which PHF6 and PRC2 complex collaborates to maintain bivalent promoters hypomethylated. Functional studies are required to detail the role of PHF6 in maintaining H3K27me3 marks at bivalent promoters and in preventing them from DNA hypermethylation during leukemogenesis.

We demonstrate that *PHF6* alterations are conserved during the clonal evolution of the disease and might provide better clonal fitness at relapse as suggested in previous publications (28,31). It has also been suggested that PHF6 LoF is an early event during leukemogenesis (28). Nevertheless, this event is not sufficient to specifically develop T-ALL, as mice lacking PHF6 develop a heterogeneous spectrum of hemopathies, and further leukemogenic events may be required to develop T-ALL, such as homeodomain protein overexpression (30). The loss of epigenetic plasticity following PHF6 alteration and DNA hypermethylation could participate in early leukemogenesis, by slowing down the differentiation program of thymocytes as it is suggested for other cancer (55). This epigenetic state, and the LoF of PHF6 itself, could lead to genomic instability and appearance of chromosomal rearrangement responsible for ectopic expression of homeobox transcription factor such as TLXs (55,58). Importantly, we showed that TLX1 does not locate at bivalent promoters suggesting that its overexpression may not be directly responsible for the epigenetic changes depicted in this study. The strong association between *PHF6* alterations and homeodomain

oncotypes also supports oncogenic cooperation. However, further studies are needed to explore the role and chronology of *PHF6* LoF in leukemogenesis.

Finally, our results show that *PHF6* alterations are associated with better overall survival, event-free survival, and lower cumulative risk of relapse, independently of other known prognostic parameters. Moreover, despite significantly older age, patient with *PHF6*<sup>ALT</sup> T-ALL achieve complete remission significantly more often and have lower incidence of death during induction treatment. However, according to this study, one out of four *PHF6*<sup>ALT</sup> patients still experienced relapsed or refractory disease. Hence, novel treatments are still needed for those patients. *PHF6* alterations are largely conserved at relapse and are therefore potential biomarkers for innovative targeted therapies. This study reveals increased in vitro and in vivo sensitivity of *PHF6*<sup>ALT</sup> T-ALL to the demethylating agent 5-azacytidine alone and in combination with the BCL2 inhibitor venetoclax. Such treatment combination has recently become a standard of care for patients diagnosed with AML and not eligible for intensive chemotherapy induction, and has been given to patients presenting R/R T-ALL (44,45). These observations led us to treat three patients bearing *PHF6* alterations with good response and minimal toxicity. Altogether, this study provides clinical evidence that *PHF6* alterations is associated with 5-azacytidine and venetoclax combination response and might therefore be used as surrogate of eligibility to this treatment.

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## Tables

Clinical characteristics						
	All N = 216	PHF6 <sup>ALT</sup> N = 102	PHF6 <sup>WT</sup> N = 114	p		
<b>Patients characteristics</b>						
Median age (CI)	30.49 (16.27 - 59.15)	34.17 (16.42 - 58.05)	28.22 (16.27 - 59.15)	0.04		
White cell count in G/L (CI)	35.15 (1.00 - 645.00)	33.00 (1.00 - 604.40)	40.45 (2.20 - 645.00)	0.90		
CNS involvement (%)	24 (11%)	10 (10%)	14 (12%)	0.67		
<b>Treatment response</b>						
Steroid response (%)	117 (54%)	59 (58%)	58 (51%)	0.34		
Early BM blast clearance (%)	119 (55%)	56 (55%)	63 (55%)	0.89		
Induction mortality rate (%)	9 (4%)	1 (1%)	8 (7%)	0.04		
Complete remission rate (%)	199 (92%)	98 (96%)	101 (89%)	0.05		
MRD1 < 10 <sup>-4</sup> (%) (N = 122)	38 (31%)	22 (33%)	16 (29%)	0.70		
HSCT (%)	77 (36%)	40 (39%)	37 (32%)	0.32		
<b>NOTCH1/FBXW7 - RAS/PTEN classifier</b>						
High risk (%)	88 (41%)	28 (27%)	60 (53%)	< 0.001		
<b>Outcomes</b>						
<b>Outcome estimations at 5 yrs</b>						
Cumulative incidence of relapse (CI)	0.33 (0.27 - 0.40)	0.27 (0.19 - 0.37)	0.39 (0.30 - 0.49)	0.04		
Event-free survival (CI)	0.56 (0.49 - 0.62)	0.63 (0.53 - 0.72)	0.49 (0.40 - 0.58)	0.02		
Overall survival (CI)	0.65 (0.58 - 0.71)	0.74 (0.64 - 0.81)	0.57 (0.47 - 0.66)	0.01		
<b>Overall survival</b>						
	<b>Univariate</b>			<b>Multivariate</b>		
	<b>HR</b>	<b>CI</b>	<b>p</b>	<b>HR</b>	<b>CI</b>	<b>p</b>
Age	1.03	1.01 - 1.05	0.002	1.05	1.03 - 1.07	<0.001
White cells blood count	1.64	1.11 - 2.40	0.01	1.94	1.29 - 2.91	0.001
CNS involvement	2.43	1.38 - 4.27	0.002	2.27	1.28 - 4.03	0.005
Steroid response	0.62	0.41 - 0.98	0.04	0.92	0.57 - 1.47	0.72
Classifier high risk	2.74	1.75 - 4.29	<0.001	2.48	1.53 - 4.02	<0.001
PHF6 <sup>ALT</sup>	0.51	0.32 - 0.80	0.004	0.58	0.36 - 0.94	0.03

**Table 1: Patients clinical characteristics and outcomes.**

CI, confidence interval; CNS, central nervous system; HSCT, hematopoietic stem cell transplant; HR, hazard ratio; CI, confidence interval; CNS, central nervous system.

## Figure legends

### Figure 1: *PHF6* alterations in the GRAALL-2003-2005 studies.

(A) Pie chart indicating the incidence of *PHF6* alterations. (B) Pie chart indicating the proportion of the different types of mutations affecting the *PHF6* gene in our cohort. (C) Lollipop plots indicating the observed mutations and their consequences. The red bars map the observed chromosomal deletions involving the *PHF6* gene in the GRAALL-2003-2005 studies. (D) Oncoplot depicting the genetic anomalies observed in *PHF6* WT or altered T-ALL cases of the GRAALL 2003-2005 studies. For each case, their immunophenotype, and ETP classification. As ETP status is separate from the immunophenotypes, some T-ALL cases are both considered as ETP and immature or TCRAB in the analyses. Genes are classified by functional groups. (E) Panel indicating alterations with significantly different frequency between *PHF6* WT or altered patients. Statistical differences were compared by Fisher tests. (F) Bar graphs showing the significant association between *PHF6* status and the maturation arrest stage. Annotations indicate the incidence of each maturation arrest stage among the entire cohort. *PHF6* WT ETP cases are also considered as immature for 16/24, cortical for 4/54, TCRGD for 1/10 and TCRAB for 1/17 cases. For the *PHF6* ALT, ETP cases count as immature for 18/28, cortical for 5/53, TCRGD for 1/13. (G) Bar graphs displaying significant association between *PHF6* status and the defining T-ALL genetic events.

### Figure 2. Clinical impact of *PHF6* alterations in the GRAALL-2003-2005 studies.

(A) Overall survival, (B) event-free survival and (C) cumulative incidence of relapse in the GRAALL 2003-2005 studies. Red curves represent the *PHF6*<sup>ALT</sup>T-ALL, and blue curves the WT patients.

**Figure 3: PHF6ALTT-ALL display highly methylated methylome.**

(A) Genome-wide average CpG methylation of PHF6WT and PHF6ALT T-ALL. (B) Number of hyper- and hypomethylated DMPs in PHF6ALTT-ALL. (C and D) Stacked barplots depicting the relative frequency of significant hyper- or hypomethylated CpGs in relation to their CpG location (C) or context (D). (E) Average CpG methylation of hyDMPs of thymocytes, PHF6WT and PHF6ALT T-ALL. (F) Number of hyper- and hypomethylated promoters in PHF6ALT T-ALL. (G) Barplot for the enrichment of hyper- and hypomethylated DMPs in transcription factors target genes and histone marks. Bar color intensity is coded for odds ratio. (H) Heatmap of H3K27me3 (left) and H3K4me3 (right) histone marks peaks at promoters of genes associated to hyDMPs or to random promoters, at different stages of thymopoiesis (ETP = early T precursors, EC = early cortical, LC = late corticals, SP = single positives) Genes are ordered identically for both marks. (I) Genome-wide average CpG methylation of PHF6WT/PRC2WT, PHF6ALT/PRC2WT, PRC2ALT/PHF6WT, PHF6ALT/PRC2ALT T-ALL. (J) Venn diagram of hyDMPs between PHF6ALT/PRC2WT, PRC2ALT/PHF6WT, PHF6ALT/PRC2ALT T-ALL versus None T-ALL. (K) Uniform Manifold Approximation and Projection of samples according to genome-wide methylation levels ( $\beta$ -values). Each dot represent a patient sample and is colored according to PHF6/PRC2 mutational status. (L) Dot plot for the enrichment of hyper-DMPs common or unique to PHF6ALT/PRC2WT, PRC2ALT/PHF6WT, PHF6ALT/PRC2ALT versus PHF6WT/PRC2WT T-ALLs, in transcription factors target genes, histone marks and gene ontology. Dots are sized-coded for adjusted p-values in the  $-\log_{10}$  scale, and color intensity is scaled according to odds ratio.

**Figure 4: PHF6ALT T-ALL display altered epigenetics landscape of bivalent promoters**

(A) Dot plot for the enrichment of TLX1 (HPB-ALL), PHF6 and TAL1 (Jurkat) peaks along with hyDMPs in active or bivalent promoters (x axis) at specific stages of thymopoiesis. Circles are color-coded for significance, dots color gradient represent odds ratio and the size of the dots represents P values in the log10 scale. (B) Heatmap representing the hierarchical clustering of T-ALL samples and normal thymocytes (x-axis) according to the 10 000 most variable CpG  $\beta$ -values associated to active (left) and bivalent (right) promoters (y-axis). (C) Uniform Manifold Approximation and Projection of T-ALL samples and normal thymocytes according to  $\beta$ -values of CpG associated to thymic bivalent promoters. Each dot represent a sample and is colored according to PHF6/PRC2 mutational status. (D) Heatmap of H3K27me3 (top) and H3K4me3 (bottom) histone marks peaks at thymic bivalent promoters in PHF6WT/PRC2WT (lightblue), PHF6ALT/PRC2WT (red), PHF6WT/PRC2ALT (yellow), PHF6ALT/PRC2ALT (black) T-ALL. (E) Venn diagram representing the proportion of thymic bivalent genes associated to hyDMPs. (F) Volcano plot of differentially expressed genes in PHF6ALT versus PHF6WT. Gene list with an absolute log fold change value superior to 2.

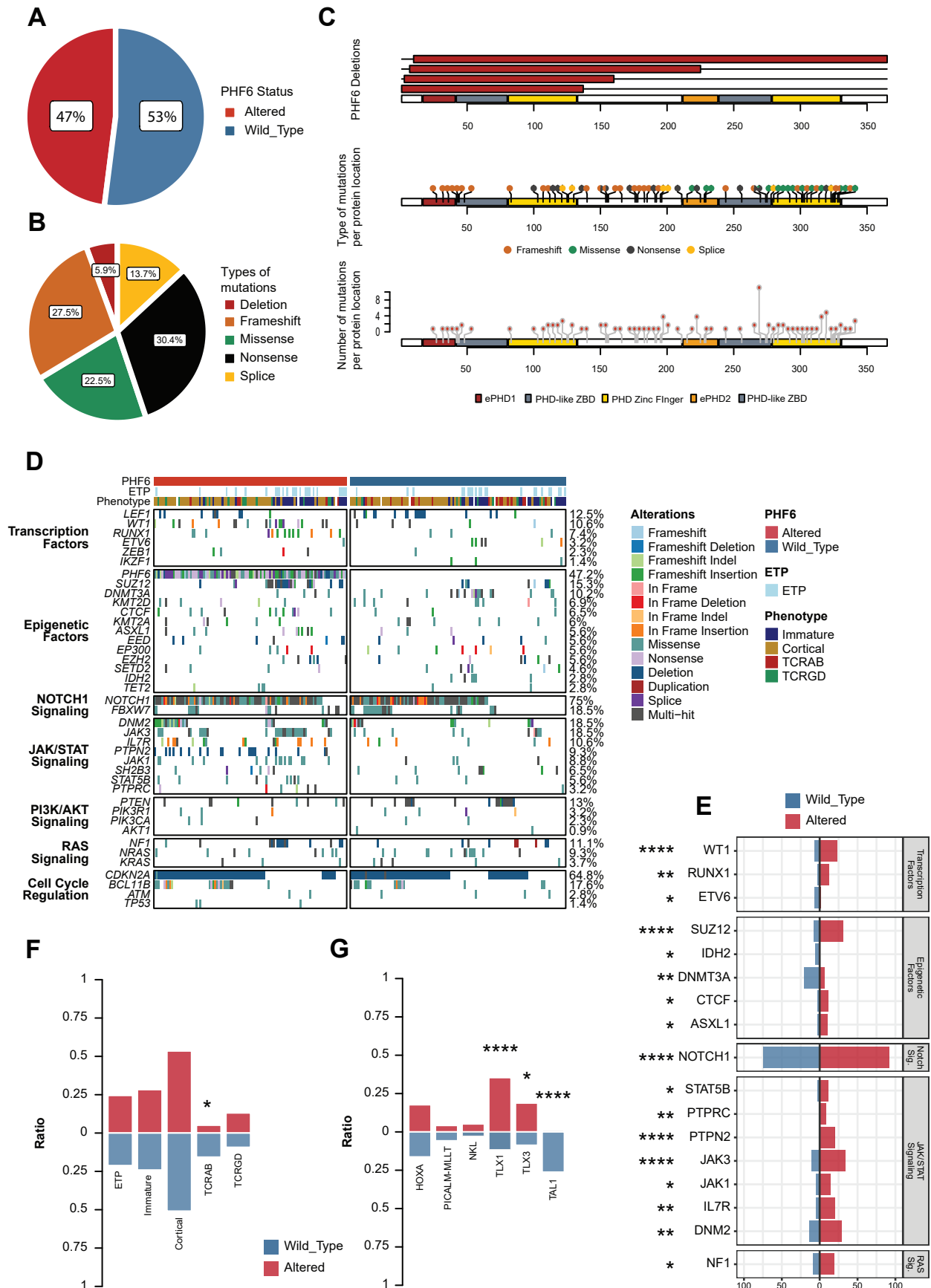
**Figure 5: 5-azacytidine and venetoclax sensitivity of PHF6<sup>ALT</sup>T-ALL**

(A) Survival for PDX mouse models of primary T-ALL : PHF6<sup>WT</sup> (UPNT-M894 : Untreated N = 5, Azacitidine N = 3 ; UPNT-M894 : Untreated N = 5, Azacitidine N = 4) and PHF6<sup>ALT</sup> ((UPNT-M640 : Untreated N = 4, Azacitidine N = 4 ; UPNT-M149 : Untreated N = 5, Azacitidine N = 5)). When blasts count reached 1% in blood, mice were administered with NaCl (Untreated : orange line) or treated with 5-azacytidine in a curative-like manner for 2 weeks (5 mg/kg per day for 5 days a week) (5-azacytidine

in a curative-like manner : green line). P values are calculated by comparing the untreated group with treated group using log-rank test. (B) Viability of T-ALL fresh samples blasts measured after 3 days of treatment with 5-azacytidine at the indicated doses. Means and SEM are plotted (C) BCL2 gene expression levels in PHF6<sup>WT</sup>/PRC2<sup>WT</sup> (lightblue), PHF6<sup>ALT</sup>/PRC2<sup>WT</sup> (red), PHF6<sup>WT</sup>/PRC2<sup>ALT</sup> (yellow), PHF6<sup>ALT</sup>/PRC2<sup>ALT</sup> (black) T-ALL (D) Viability of T-ALL fresh samples blasts measured after 3 days of treatment with 5-azacytidine at the indicated doses combined with 125  $\mu$ M of Venetoclax. Means and SEM are plotted. (E) Clinical timelines of three patient diagnosed with R/R T-ALL treated with 5-Azacytidine and Venetoclax. Timelines are centered on the time of initiation of the combination.

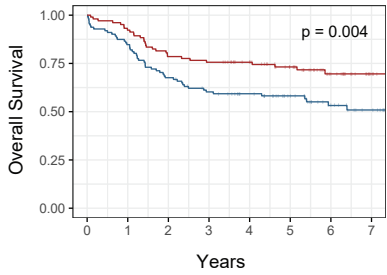


# Figure 1

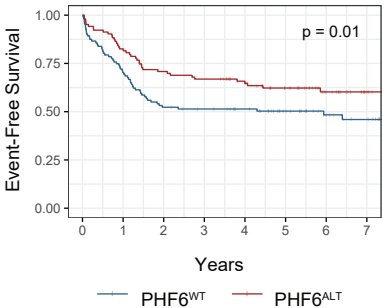


**Figure 2**

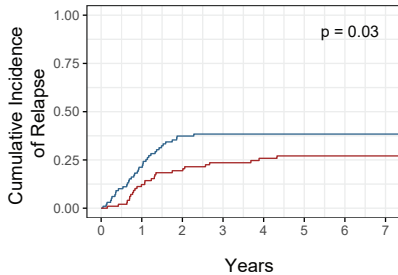
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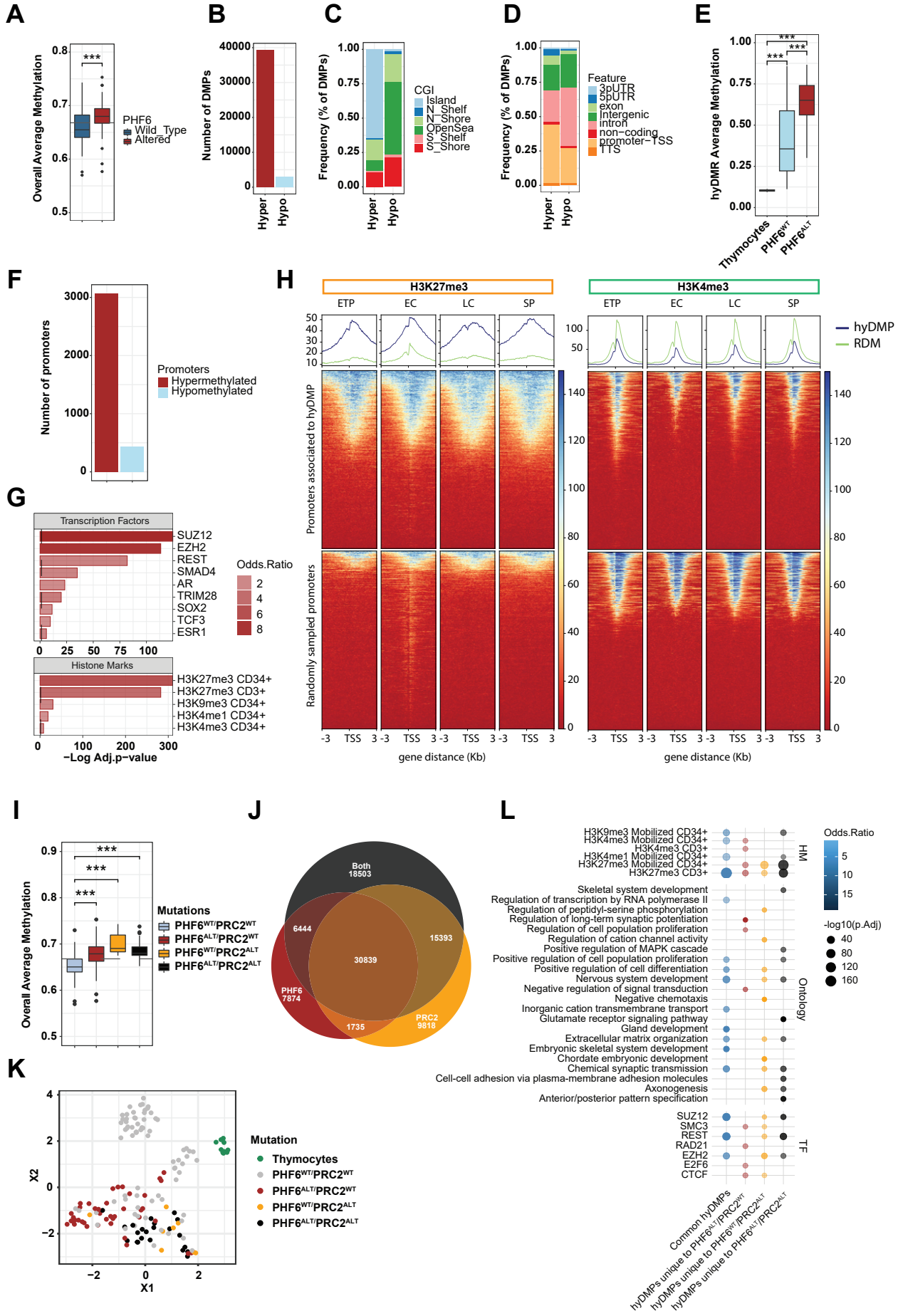
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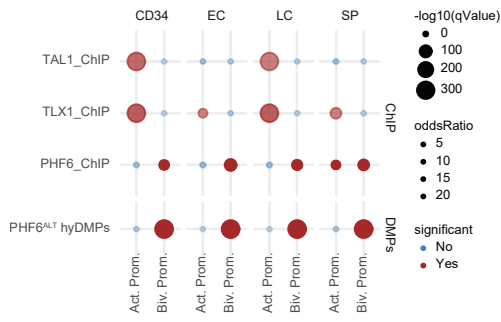


# Figure 3

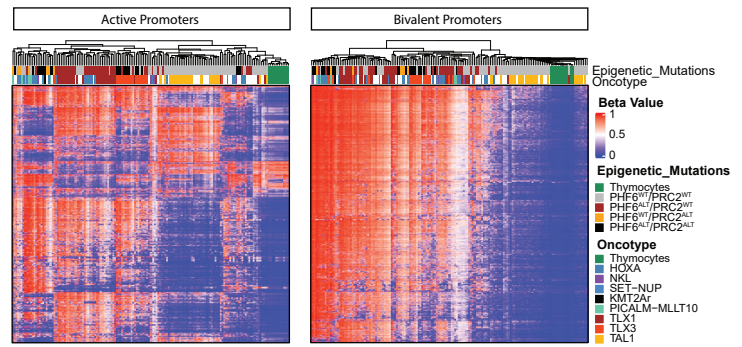


# Figure 4

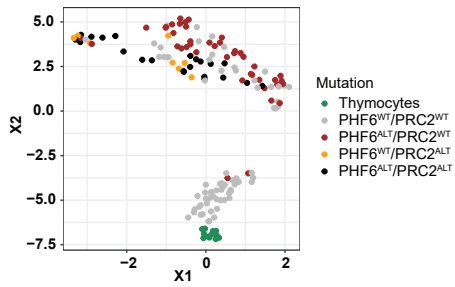
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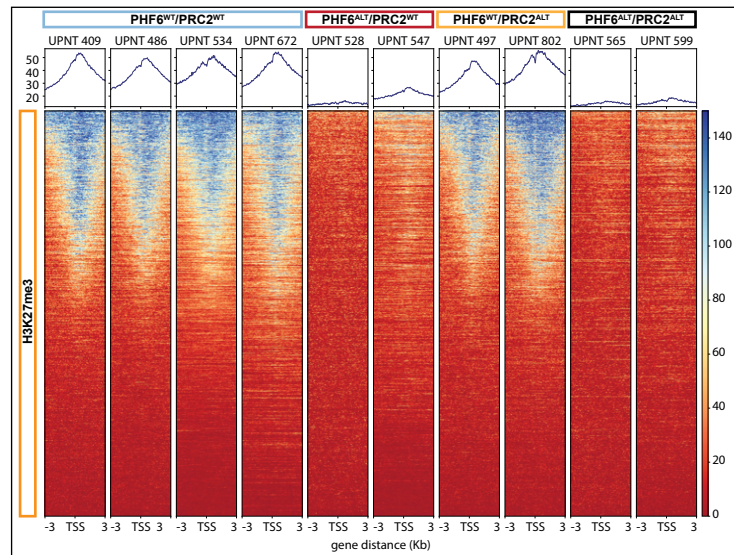
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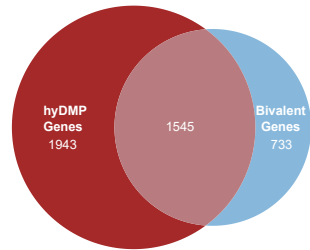
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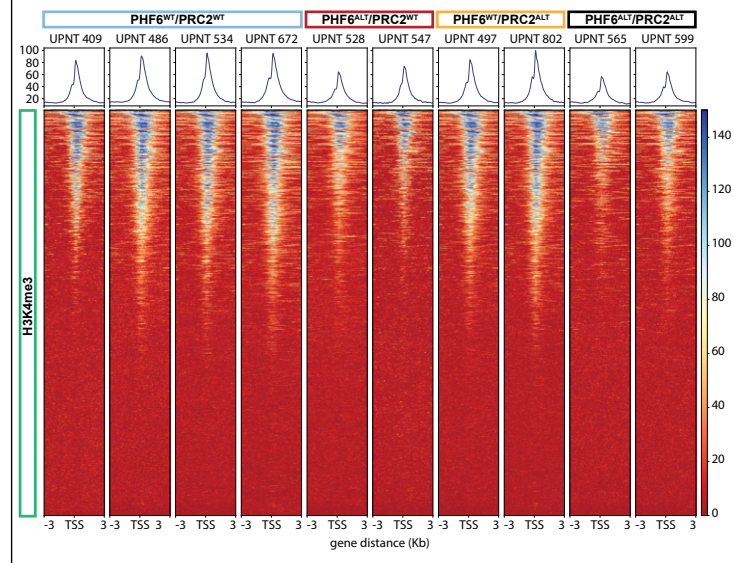
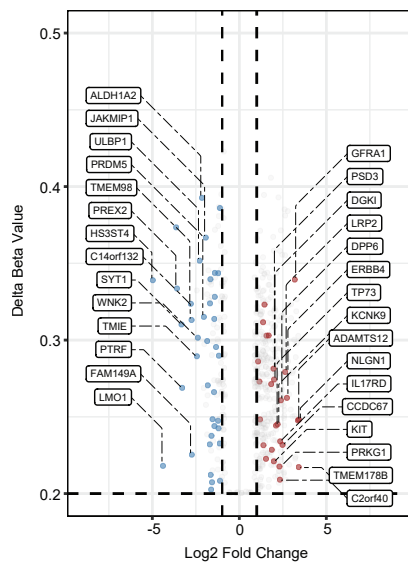
**D**



**E**

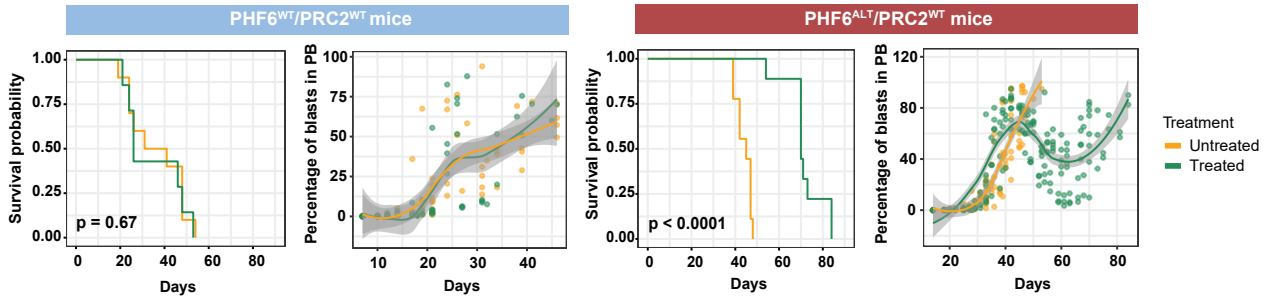


**F**

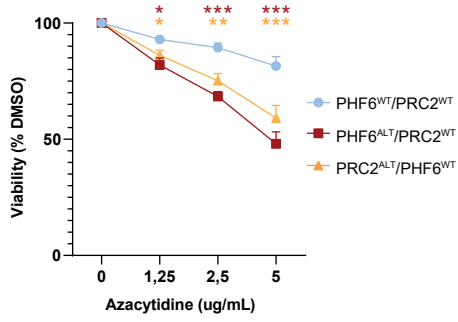


**Figure 5**

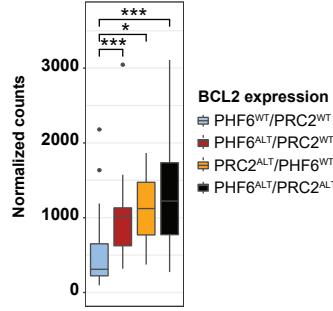
**A**



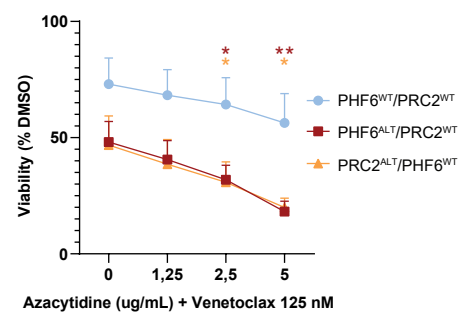
**B**



**C**



**D**



**E**

