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**Article**

**Hif-1α Deletion May Lead to Adverse Treatment Effect in a Mouse Model of MLL-AF9-Driven AML**

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**SUMMARY**

Relapse of acute myeloid leukemia (AML) remains a significant clinical challenge due to limited therapeutic options and poor prognosis. Leukemic stem cells (LSCs) are the cellular units responsible for relapse in AML, and strategies that target LSCs are thus critical. One proposed potential strategy to this end is to break the quiescent state of LSCs, thereby sensitizing LSCs to conventional cytostatics. The hypoxia-inducible factor (HIF) pathway is a main driver of cellular quiescence and a potential therapeutic target, with precedence from both solid cancers and leukemias. Here, we used a conditional knockout Hif-1α mouse model together with a standard chemotherapy regimen to evaluate LSC targeting in AML. Contrary to expectation, our studies revealed that Hif-1α-deleted-leukemias displayed a faster disease progression after chemotherapy. Our studies thereby challenge the general notion of cancer stem cell sensitization by inhibition of the HIF pathway, and warrant caution when applying HIF inhibition in combination with chemotherapy in AML.

**INTRODUCTION**

Acute myeloid leukemia (AML) is a clonal disorder characterized by a rapid accumulation of differentiation-arrested myeloid blasts. Remission can be achieved in most patients by a combination of cytarabine (AraC) and anthracycline therapy. However, few treatment improvements have been observed over the past decades (Rowe and Tallman, 2010) and 40%–60% of patients still relapse (Burnett et al., 2011; Yanada et al., 2008), which is associated in general with limited treatment options (Dohner et al., 2010).

Relapse is attributed to a minor subpopulation of cells referred to as leukemic stem cells (LSCs). LSCs are resilient to cytotoxic effects of chemotherapy, via mechanisms that include resistance to apoptosis, increased capacities to efflux drugs, and relative quiescence (Thomas and Majeti, 2017). This quiescence in turn underlies resistance to compounds that target energy metabolism (Essers and Troupp, 2010).

Cellular quiescence is typically related to activation of hypoxia-signaling pathway, driven by hypoxia-inducible factor (HIF) complex, which constitutes a family of 3 heterodimeric transcription factors HIF-1, HIF-2, and HIF-3 (Flamme et al., 1997; Makino et al., 2002; Semenza and Wang, 1992). The oxygen-dependent regulation of HIFs depends on stabilization of an associated α subunit. Hence, at oxygen levels above 5%, the α subunit is proteasomally degraded. In contrast, under hypoxic conditions the α subunit is stabilized, dimerizes with constitutively expressed β subunit, and promotes transcription of target genes regulating key cellular processes such as angiogenesis, proliferation, metabolism, and apoptosis (Semenza, 2003).

The most primitive hematopoietic stem cells (HSCs) reside in hypoxic niches in the bone marrow (BM) (Morisson and Scadden, 2014). Thereby, HSCs are kept dormant and free of genetic changes that mostly occur during DNA replication. However, while HIF-1α signaling was reported to critically regulate HSC maintenance (Takubo et al., 2010), other studies showed that depletion of HIF-1α/HIF-2α had no such effect (Guitart et al., 2013). The HIF pathway has also been proposed to play important roles in AML, a notion supported by the preferential localization of chemotherapy-resistant AML cells to the hypoxic endosteal niche in the BM (Ishikawa et al., 2007). Accordingly, several studies have shown that loss of HIF-1/2 leads to abrogation of LSCs (Rouault-Pierre et al., 2013; Wang et al., 2011; Zhang et al., 2012). Contrasting these results are studies demonstrating that loss of HIF-1/2 does not impact on mouse models of AML, or alternatively can give rise to an even more severe leukemic phenotype (Velasco-Hernandez et al., 2014; Vukovic et al., 2015). Nonetheless, targeting hypoxia and HIFs has been considered a key anticancer approach (Frolova et al., 2012; Rouault-Pierre et al., 2013; Wang et al., 2011).

Here, we investigated the possibility to sensitize LSCs from a mouse model of AML to a clinically relevant chemotherapy regimen by targeting the hypoxia pathway via Hif-1α. Contrary to expectation, our work suggests a more...
aggressive disease outcome upon Hif-1α deletion, which challenges the general use of hypoxia targeting for therapeutic benefit in AML.

RESULTS

Deletion of Hif-1α Accelerates the Progression of Chemotherapy-Treated Leukemia

To study effects of Hif-1α on the sensitivity of LSCs to chemotherapy, we bred Hif-1αβ mice (Ryan et al., 2000) with Rosa26Cre-ER12 mice (Ventura et al., 2007), generating a model that allows for Hif-1α deletion upon tamoxifen treatment (Figure S1A). c-Kit+ cells from Hif-1αβ;Rosa26Cre-ER12/+ mice were transduced with MLL-AF9-GFP retrovirus and transplanted into mice, which led to development of MLL-AF9-driven AML. Primary leukemias were next used to study AML progression in secondary hosts.

After comparing two different chemotherapy protocols (Figure S1B), we used the one described by Zuber et al. (2009), which resulted in severely decreased white blood cells, erythrocytes, and platelets, akin to what is observed in human patients. We then titrated the latency of the disease using different leukemic cell doses (Figure S1C) and chose a dose of 10^4 cells, which ensured a sufficiently long latency for posterior procedures. Cells from individual MLL-AF9-leukemic mice were transplanted into new recipients (Figure 1A). As expected, disease progressed more slowly in the chemotherapy-treated (CTX) mice (Figures 1B–1F). More intriguingly, we observed a faster disease evolution upon Hif-1α deletion (Figures 1B–1F and S2). As expected, disease progressed more slowly in the chemotherapy-treated (CTX) mice (Figures 1B–1F). More intriguingly, we observed a faster disease evolution upon Hif-1α deletion. This was seen in both CTX and PBS groups (Figures 1C–1F). When we assessed phenotypically more primitive GFP− c-Kit+ or GFP− Lin− c-Kit+ cells, we observed a higher abundance in the Hif-1α-deleted (Hif-1α−/−) than in the Hif-1α-intact (Hif-1α+/+ or Hif-1α−/+) group after chemotherapy (Figures 1G, 1H, and S2). While these populations were very infrequent in BM of PBS groups, with no clear differences between Hif-1α−/− and Hif-1α+/+ cells, a higher number of such cells upon Hif-1α deletion were observed in the spleens of the same animals. Taken together, these results demonstrate an increment of potential LSCs (c-Kit+) upon Hif-1α−/− deletion, which was particularly significant after chemotherapy.

Deletion of Hif-1α Does Not Decrease the Frequency of LSCs after Chemotherapy Treatment

To evaluate the impact of chemotherapy and Hif-1α deletion on LSCs more directly, we performed limiting dilution analysis (LDA) (Figure 2A).

For an in vitro approach, we sorted 4, 8, or 12 GFP+ leukemic BM cells from each group into individual wells. Proliferation was next evaluated in both normoxic (20% oxygen) and hypoxic (1% oxygen) conditions (Figures 2B and 2E). We obtained the highest LSC frequency from PBS-Hif-1α−/− cells (1/12.3 and 1/21.7 in hypoxia or normoxia, respectively, means from 3 independent experiments). Within CTX-treated cells, Hif-1α−/− samples contained a higher LSC frequency compared with Hif-1α+/+ cells (1/26.8 versus 1/47.1 in hypoxia, and 1/40.4 versus 1/94.4 in normoxia, respectively, means from 3 independent experiments), although this failed to reach statistical significance.

We next assessed LSC frequencies in vivo, injecting 10, 10^2, 10^3, or 10^4 GFP+ BM cells from each group into new recipients (Figure 2C). While transplantation of 10^3 and 10^4 cells resulted in 100% mortality, lower doses of 10 and 10^2 cells indicated an LSC frequency highest in the PBS-Hif-1α−/− group (1/18.8). Among CTX-treated samples, we observed a higher LSC frequency in the Hif-1α−/− group (1/34.5) than in the Hif-1α+/+ group (1/56.1) (Figures 2D and 2E).

Estimation of total number of LSCs in the BM (Figure 2F) demonstrated that deletion of Hif-1α fails to reduce LSC frequencies/numbers after chemotherapy.

Hif-1α Deletion Affects Transcriptional Expression of Replication, Transcription, and Translation-Related Genes

Our data indicated that Hif-1α deletion contributed to a more rapid disease progression (Figure 1). To tease out a potential mechanism, we therefore conducted single-cell RNA sequencing on leukemic cells from the 4 evaluated settings (Hif-1α−/−/Hif-1α+/+ × chemotherapy) (Figure 3A). Aggregated data were subjected to dimensionality reduction using the t-distributed stochastic neighbor embedding (t-SNE) method (Figure 3B), revealing dramatically different transcriptional profiles of cells following chemotherapy, but less different according to Hif-1α status. This indicates either a modification of transcription upon chemotherapy or, more likely, a selective survival regardless of Hif-1α status.

By K-means clustering, we divided the entire dataset into 10 different clusters. We observed that clusters with functions related to more differentiated cells (e.g., immune system process, response to interleukin-1/tumor necrosis factor, innate immunity or response to virus/interferon) were predominant in PBS samples, while
others (e.g., mitotic cell cycle, cellular response to DNA damage, and transcriptional regulation) were associated with CTX samples. Multiple genes related to myeloid differentiation were predominantly expressed in PBS samples (data not shown), strongly suggesting effective chemotherapy targeting of more mature leukemia-associated cells.

Pairwise sample comparisons produced a collection of 331 differentially expressed genes (Figures S3 and 3C). When comparing significantly overexpressed genes among groups (adjusted p value <0.05), we found a very different distribution of cells between CTX and PBS samples, but only a small subset of genes uniquely expressed in either Hif-1α−/− or Hif-1α+/+ cells (Figures 3D and S4). In
**Figure 2. Deletion of Hif-1α Does Not Decrease LSC Frequency after Chemotherapy**

(A) Experimental design for the analysis of LSC frequency.

(B) Limiting dilution analysis (LDA) from *in vitro* data. Variable numbers of GFP+ cells (4, 8, or 12 cells) from the 3 experimental groups were cultured *in vitro* under normoxic or hypoxic conditions and proliferative capacity was evaluated after 10 days. LSC frequencies were calculated using ELDA software. Plots show 1 representative of 3 independent experiments (cells from 3 different donors) with 96 replicates per group and condition.

(C) Survival curves of mice transplanted with 10 or 100 GFP+ cells from mice treated with the combination of tamoxifen + chemotherapy (KO-CTX), tamoxifen + PBS (KO-PBS), or oil + chemotherapy (wt-CTX) (n = 4–6 mice per group from 1 experiment).

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single-cell analysis, this indicates that there are differences in transcriptional regulation or that analyzed cells from each sample are different.

To identify functional protein association networks among differentially expressed genes, we used the STRING database (Figures 3E and 4). For the 4 sets of differentially expressed genes (log fold change >0.4, adjusted p value <0.05), we obtained a significant protein-protein interaction (PPI) enrichment score, indicating that there are more interactions among these proteins than a random set of proteins from the genome (PPI enrichment p values: Hif-1α+/−-CTX = 0; Hif-1α+/−-CTX = 1.03 × 10−6; Hif-1α−/−-PBS = 0; Hif-1α−/−-PBS = 0.0251). This indicates that genes in each set are at least partially biologically connected. To correlate these genes with a biological function, we performed gene ontology and KEGG pathway enrichment analyses (Figure 3E). This revealed that mRNA processing, DNA replication, and protein folding were significantly overrepresented in the Hif-1α−/−-CTX group, while the Hif-1α+/−-CTX group was associated with the HIF-1α pathway, glycolysis, and other carbon metabolism processes. In PBS groups, translation was the most represented pathway associated with Hif-1α−/− cells, while no significant pathways could be assigned to Hif-1α+/+ cells.

**DISCUSSION**

Relapse of AML remains a major therapeutic challenge, not least because of the few therapeutic options for many of these patients, which in most cases is restricted to palliative care.

One potential strategy for eradicating LSCs, responsible for relapse in AML, is to sensitize them to chemotherapy. We here explored whether targeting the HIF pathway could be used for this using an MLL-AF9 murine AML model, which was subjected to a standard and clinically relevant chemotherapy protocol. In our work, Hif-1α−/− cells displayed a faster evolution of the disease compared with Hif-1α+/+ cells following chemotherapy. In our work, Hif-1α−/− cells were critical for LSCs maintenance. In our work, we used the Rosa26Cre-ERT2 rather than the more commonly used Mx1-Cre model, which is associated with significant leakiness (Velasco-Hernandez et al., 2016). This allowed us to study the effect of Hif-1α inhibition specifically in AML cells of the same founder clone, which circumvents the risk that different leukemic clones have intrinsically different behavior. Inducing the deletion in Rosa26Cre-ERT2 mice requires injection of tamoxifen that might itself affect the initiation of MLL-AF9 leukemia (Sanchez-Aguilera et al., 2014). However, as the described effect of tamoxifen is a regression of the disease, the isolated effect of knocking out Hif-1α could be even more pronounced than the one we report.

The standard chemotherapy regimen for AML is a combination of AraC and an anthracycline (such as doxorubicin). Doxorubicin has itself been attributed Hif-inhibiting properties (Lee et al., 2009). Different disease progression in CTX-Hif-1α−/− and Hif-1α+/+ samples indicates that even if doxorubicin is inhibiting HIF function, this effect is not complete and can be further increased by alternative means.

There is evidence that c-Kit deregulation and overexpression could be factors contributing to the chemotherapy resistance in AML (Advani et al., 2008). The higher numbers of c-Kit+ cells in the Hif-1α−/− chemotherapy group might therefore be an indicator of higher drug resistance of these cells, with c-Kit being a well-described stem cell marker of murine MLL-AF9-driven LSCs (Krivtsov et al., 2006). However, our c-Kit expression data correlated poorly with LSC frequencies measured by LDA. Previously it was thought that chemotherapy promoted LSC enrichment (Ishikawa et al., 2007; Saito et al., 2010). However, although cytarabine decreases the frequency of LSCs in human AML models (Griessinger et al., 2014), it fails to promote enrichment either of quiescent cells or more primitive CD34+/CD38− cells, emphasizing that functional assessments are critical. We did not observe an increment in LSC frequencies measured by LDA but observed a substantial reduction (around 3-fold) of the total bulk of leukemic cells, indicating an overall enrichment of LSCs following chemotherapy. In addition, we observed an increased frequency of c-Kit+ cells by Hif-1α deletion, which further supports the increased aggressive phenotype of Hif-1α−/− cells.

Hif-1α is rapidly degraded in normoxia and Hif-1α+/+ cells should thus theoretically behave as Hif-1α+/− cells, which
Figure 3. Single-Cell RNA Sequencing of Leukemic Cells after Chemotherapy Treatment

(A) Experimental design for single-cell RNA-sequencing analysis. After treatment with tamoxifen/oil and chemotherapy/PBS, GFP+ cells were collected at day 28 after transplantation and subjected to RNA processing.

(B) t-SNE plots representing gene expression profiles of all individual cells analyzed: KO-CTX (n = 1,217 cells), KO-PBS (n = 1,696 cells), wt-CTX (n = 1,301 cells), and wt-PBS (n = 1,450 cells). Each dot represents one cell. K-means clustering groups cells into 10 clusters, which are represented by different colors. Differentially expressed genes in each cluster were correlated with the main biological function indicated at the right side of the plots.

(C) Heatmap depicting significantly differentially expressed genes in single cells. Heatmap with the full set of genes can be found in Figure S3. Hif-1α target genes are indicated by black dots.

(D) Venn diagrams showing the distribution among groups of differentially expressed genes (adjusted p value <0.05). Of note, there are no overlapping genes between chemotherapy- and PBS-treated groups among this set of differentially expressed genes.

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would suggest similar LSC frequencies between Hif-1α−/− and Hif-1α+/+ cells in normoxia. Somewhat surprisingly, we observed a higher LSC frequency of Hif-1α−/− cells, although not significant (p = 0.1079). One possibility is that this indicates the involvement of one or more Hif-1α-oxygen-independent mechanisms. For instance, it is well established that Hif-1α can be induced by the PTEN/PI3K/AKT axis (Zundel et al., 2000), a pathway that appears to extend also to MLL-AF9 leukemia (Hoshii et al., 2012).

After chemotherapy, we observed a more aggressive phenotype as a consequence of Hif-1α deletion. We hypothesized that this faster development of disease could

![Figure 4. Protein Association Networks Affected by Chemotherapy](image)

Figure 4. Protein Association Networks Affected by Chemotherapy

Protein association network analysis of differentially expressed set of genes (average log fold change >0.4, adjusted p value <0.05). Uniquely expressed genes from each group are indicated by its assigned color. Common genes among KO and wt samples, within each specific set (CTX- or PBS-treated), are shown in dark gray. Associated biological function to each gene is indicated, with the color of the border showing the main processes regulated by these genes. Interaction enrichment score >0.5. CTX, chemotherapy.

(E) Gene ontology (GO) and KEGG pathway enrichment analysis from uniquely differentially expressed genes in each group indicating the associated biological process. CTX, chemotherapy; IL1/TNF, interleukin-1/tumor necrosis factor; INF, interferon; dim, dimension; FDR, false discovery rate. See also Figures S3 and S4.
be explained by two scenarios: (1) leukemic Hif-1α−/− cells are less sensitive to the chemotherapy, which would be in contrast to our original expectation; and/or (2) leukemic Hif-1α−/− cells expand faster than Hif-1α+/+ cells after chemotherapy. To discriminate between these two options and unveil possible mechanisms leading to this effect, we conducted a single-cell transcriptome analysis.

We found that chemotherapy-treated cells displayed a distinct transcriptomic profile when compared with PBS-treated cells, but with fewer differences when comparing Hif-1α−/− and Hif-1α+/+ cells. Still, of the significantly over-expressed genes specific for each group, we identified an increment in genes involved in translation (PBS-Hif-1α−/− cells) and in mRNA processing, DNA replication, and protein folding (CTX-Hif-1α−/− cells). Overall, this is in line with a more active proliferative status of Hif-1α−/− cells and suggests that deletion of Hif-1α is transcriptionally deregulating these aforementioned specific pathways. Chemotherapy-depleted cells with an immune-related phenotype represent the cells expanded in leukemia. As expected, Hif-1α−/− cells were depleted in expression of Hif-1α targets, and Hif-1α deletion leads to an enhanced transcription of genes involved mainly in replication, mRNA processing, and translation.

These results are in accordance with the described effect of Hif-1α, inhibiting cell cycle and promoting quiescence, actions attributed to blockage of c-MYC and enhancement of cyclin-dependent kinase inhibitors such as p21/CDKN1A (Koshiji et al., 2004). Since we are deleting only one member of a multigene family, it remains a possibility that other members, i.e., Hif-2α, could have a compensatory effect. Future work could benefit from an increased understanding of both gene redundancy and potential hypoxia-independent effects of Hif-1α.

Overall, our data suggest that Hif-1α inhibition failed to improve the outcome of chemotherapy in MLL-AF9-driven AML and, in contrast, led to a faster progression of disease upon withdrawal of the treatment. Further investigations are needed to extend these conclusions to other genetic subtypes of AML.

Deletion of Hif-1α in our setting was permanent, which would be a different scenario than in a clinical setting where administration of Hif-1α inhibitors would be transient. Our results nevertheless emphasize that the effects of HIF inhibition have to be further investigated before this strategy can be applied in a clinical setting.

EXPERIMENTAL PROCEDURES

Mice
Hif-1αlox mice (Ryan et al., 2000) (JAX 007561) were crossed with the tamoxifen-inducible Rosa26Cre-ERT2 mice (Ventura et al., 2007) (JAX 008463) to generate a combined conditional knockout (KO) model. Mice were maintained at the animal facility of the Biomedical Center at Lund University (Sweden) and all experiments were performed with consent from a local ethics committee.

Statistical Analysis
All data are expressed as the mean ± SEM. Differences between groups were assessed using unpaired Student’s t tests. All analyses were performed with Prism software, version 7.0 (GraphPad Software).

Detailed descriptions of experiments can be found in Supplemental Experimental Procedures.

ACCESSION NUMBERS
The accession number for transcriptomic data is GEO: GSE119484.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures and four figures and can be found with this article online at https://doi.org/10.1016/j.stemcr.2018.11.023.

AUTHOR CONTRIBUTIONS
T.V.-H. designed the study, performed experiments, analyzed data, and wrote the manuscript; S.S. performed bioinformatics analysis; I.H. and E.E. performed experiments; and J.C and D.B. designed the study and wrote the manuscript. All authors revised the manuscript and approved the final version of this study.

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