Article

Activation of Vitamin D Receptor Pathway Enhances Differentiating Capacity in Acute Myeloid Leukemia with Isocitrate Dehydrogenase Mutations

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Simple Summary: Around 15% of acute myeloid leukemia (AML) patients harbor mutations in isocitrate dehydrogenases (IDH) which lead to the production of the oncometabolite 2-hydroxyglutarate (2-HG). Inhibitors of mutant IDH enzymes and their 2-HG production have been approved by the FDA to be used in patients. However, 60% of IDH mutant AML patients do not respond to these inhibitors or develop mechanisms of resistance, leading to relapse. Among these mechanisms, some produce a 2-HG rebound. Alternative therapies exploiting the 2-HG-dependent molecular effects could therefore be of clinical interest. In this study, we demonstrate that 2-HG specifically activates Vitamin D receptor (VDR) in IDH mutant AML cells leading to increased sensitivity to the combination of Vitamin D (or VDR agonist) and all-trans retinoic acid and revealing a new therapeutic approach that can be readily applied to AML patients in this subgroup.

Abstract: Relapses and resistance to therapeutic agents are major barriers in the treatment of acute myeloid leukemia (AML) patients. These unfavorable outcomes emphasize the need for new strategies targeting drug-resistant cells. As IDH mutations are present in the preleukemic stem cells and systematically conserved at relapse, targeting IDH mutant cells could be essential to achieve a long-term remission in the IDH mutant AML subgroup. Here, using a panel of human AML cell lines and primary AML patient specimens harboring IDH mutations, we showed that the production of an oncometabolite (R)-2-HG by IDH mutant enzymes induces vitamin D receptor-related transcriptional changes, priming these AML cells to differentiate with pharmacological doses of ATRA and/or VD. This activation occurs in a CEBPa-dependent manner. Accordingly, our findings illuminate potent and cooperative effects of IDH mutations and the vitamin D receptor pathway on differentiation in AML, revealing a novel therapeutic approach easily transferable/immediately applicable to this subgroup of AML patients.

Keywords: AML; IDH; Vitamin D; VDR; ATRA; CEBPa, Differentiation
1. Introduction

Recurrent mutations in isocitrate dehydrogenases IDH1 or IDH2 are detected in more than 15% of newly diagnosed AML patients and lead to the production of the oncometabolite (R)-2-hydroxylutarate (2-HG) [1]. IDH mutation and 2-HG induce a profound re-modeling of the epigenetic landscape of AML cells, disrupting hematopoietic differentiation [2], [3]. In this context, we have previously shown that IDH1 mutation primes AML cells to the myeloid differentiation pathway through a 2-HG-mediated activation of the transcription factor CEBPα that sensitizes IDH1 mutant (IDH1MUT) AML cells to differentiation therapy using all-trans retinoic acid (ATRA) [4]. Consistent with this study, Mugi

Here, we report that 2-HG and IDH mutation activated the vitamin D3 receptor (VDR) pathway in a CEBPα-dependent manner in AML. This favored IDH mutant AML cells to VD-induced myeloid differentiation and enhanced its synergistic effect with ATRA. This study provides a scientific rationale to treat IDH mutant AML subgroup, as an alternative to IDH-inhibitor treatment or for patients relapsing from it.

2. Materials and Methods

2.1. Primary AML cells

De-identified primary AML patient specimens are from Toulouse University Hospital (TUH) (Toulouse, France). Frozen samples were obtained from patients diagnosed with AML at TUH after signed written informed consent for research use in accordance with the Declaration of Helsinki, and stored at the HIMIP collection (BB-0033-00060). According to the French law, HIMIP biobank collection has been declared to the Ministry of Higher Education and Research (DC 2008-307, collection 1) and obtained a transfer agreement for research applications (AC 2008-129) after approbation by our institutional review board and ethics committee (Comité de Protection des Personnes Sud-Ouest et Outremer II). Clinical and biological annotations of the samples have been declared to the CNIL (Comité National Informatique et Libertés, i.e. Data processing and Liberties National Committee). Patient characteristics are summarized in Figure S1A. Human primary AML cells were grown in H4230 methylcellulose medium (STEMCELL Technologies) supplemented with 10% 5637-conditioned medium with or without ATRA (from 0.1µM to 1µM) and 1,25-Dihydroxyvitamin D3 (VD; from 10nM to 100nM) alone or in combination during 7 to 14 days at 37°C with 5%CO2.

2.2. Mice and Patient-derived xenografted model

De-identified animals were used in accordance with a protocol reviewed and approved by the Institutional Animal Care and Use Committee of Region Midi-Pyrénées (France). NSG mice were produced at the Génoult-Anexplo platform at Toulouse, France, using breeders obtained from Charles River Laboratories. Mice were housed in sterile conditions using high-efficiency particulate arrestance filtered microisolators and fed with irradiated food and sterile water.
Mice (6–9 weeks old) were sub-lethally treated with busulfan 30 mg/kg 24 h before injection of leukemic cells. Leukemia sample of IDH1 \textsuperscript{R132H} AML patient was thawed at room temperature, washed twice in PBS, and suspended in Hank’s Balanced Salt Solution at a final concentration of \(2 \times 10^6\) cells per 200 µl of Hank’s Balanced Salt Solution per mouse for tail vein injection. Daily monitoring of mice for symptoms of disease (ruffled coat, hunched back, weakness, and reduced mobility) determined the time of killing for injected animals with signs of distress. No signs of distress were seen and mice were initially analyzed for engraftment 10 wk after injection.

2.2. AML cell lines

Human AML cell lines HL60, U937 were purchased at DSMZ (Braunschweig, Germany) and KG1a at ATCC (Manassas, VA, USA). THP1 expressing IDH1 \textsuperscript{WT} were gifted by Prof. Steven Chan \cite{11}. Clones from the HL60 cell line expressing either IDH1 \textsuperscript{WT} (clone 2 and 7) or IDH1 \textsuperscript{R132H} (clone 5 and 11) were generated by our team \cite{4}. HL60 was engineered to express IDH2 \textsuperscript{WT} or IDH2 \textsuperscript{R172K} as described below. All the cells excluding THP1 were maintained in MEM\(\alpha\) with Glutamax (Gibco, Life Technologies) supplemented with 10% FBS (Invitrogen) in the presence of 100U/ml of penicillin and 100µg/ml of streptomycin. HL60 was engineered to express IDH2 \textsuperscript{WT} or IDH2 \textsuperscript{R172K} as described below. All the cells excluding THP1 were maintained in MEM\(\alpha\) with Glutamax (Gibco, Life Technologies) supplemented with 10% FBS (Invitrogen) in the presence of 100U/ml of penicillin and 100µg/ml of streptomycin. All cell lines have been routinely tested for mycoplasma contamination in the laboratory.

2.3. Reagents

Octyl-2HG (1-Octyl ester of (R)-2-HG) was synthesized at the Organic Synthesis Core (Memorial Sloane-Kettering Cancer Center, New York, NY) as previously described \cite{12} and dissolved in DMSO. Octyl-OH (1-Octanol, 8.20931, Sigma) solution in DMSO as the same concentration of octyl-2-HG was used as a control solution to assure that only the effect of (R)-2-HG-free acid on the cells was observed. ATRA (R2625, Sigma) was dissolved in DMSO.

2.4. Lentiviral infection

Lentiviral infection was carried out as previously described \cite{13}. Briefly, each construct (6µg) was co-transfected using lipofectamine 2000 with p8.1 (4µg) and pVSV-G (2µg) (providing packaging and envelope proteins) into 293T cells to produce lentiviral particles. At about 72h post transfection, 293T culture supernatants containing lentiviral particles were harvested and filtered. On the same day, HL60 or clones from HL60 cell line were infected with the lentiviral particles and polybrene (8µg/µl) by spinoculation (800g, 60min, 37\(^{\circ}\)C). After 4-6 hours, AML cells were washed with PBS1X, then diluted in classical culture media. Three days after lentiviral infection, cells were selected with hygromycin B (300µg/ml) for generation of HL60 IDH2 \textsuperscript{WT} and IDH2 \textsuperscript{R172K} or with puromycin (1µg/ml) for CEBP\(\alpha\) and CEBP\(\beta\)-LIP knockdown. For CEBP\(\alpha\) and CEBP\(\beta\)-LIP invalidation, pLKO vectors containing the following shRNA sequences were used (5’>3’): shRNA control, purchased from Sigma (SHC002 MISSION pLKO.1-puro Non-mammalian shRNA control; CCGGCAACGATGTAAGGACACCCAATCT); shRNA CEBP\(\alpha\), purchased from Sigma (SHCLNG-NM_004364, TRCN0000007306; CCGGCACAGGACGTCCTACGACATCTCGAGATGTCGATGAGCACTCTCGTGTTT); shRNA CEBP\(\beta\), purchased from Sigma (SHCLNG-NM_005194, TRCN0000007440; CCGTGTTGTATTTAAAGAA). For the generation of IDH2 \textsuperscript{WT} or IDH2 \textsuperscript{R172K} HL60, the following pSLIK vectors were used: pSLIK-IDH2-FLAG (Addgene plasmid #66806); pSLIK-IDH2-R172K-FLAG (Addgene plasmid #66807).
2.5. RNA extraction and RT-qPCR

1µg RNA was isolated using the TRI-reagent and then reverse transcribed with the iScript™ cDNA Synthesis kit. The resulting cDNAs were quantified by real-time qPCR using the SYBR Green Master mix on a Applied Biosystems StepOne™. The relative mRNA levels were calculated using the 2^(-ΔΔCT) method and normalized to PPIA (cyclophilin A) mRNA level as housekeeping gene.

<table>
<thead>
<tr>
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<td>PPIA*</td>
<td>5’TCTGAATAAGTTTGACTTGTGTTT 3’</td>
<td>5’TCTAGGCATGGGAGGGAACA 3’</td>
</tr>
<tr>
<td>RARA</td>
<td>5’GTCTTTGCGCTTGCACCAACA 3’</td>
<td>5’GCCCTCTGATTTCTCCAACA 3’</td>
</tr>
<tr>
<td>VDR</td>
<td>5’CTTCACCATGGACGACATG 3’</td>
<td>5’CTTTGGTCACGTCACT 3’</td>
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2.6. Western Blot

Proteins were analyzed by Western Blot as previously described[13]. List of antibodies used in this work: anti-CEBPα (2841, CST, 1:1000), anti-CEBPβ (sc-150, Santa Cruz, 1:1000), anti-GAPDH (5174, CST, 1:10000), anti-HSP90 (4874, CST, 1:1000), anti-IDH2 (ab55271, Abcam, 1:1000), anti-IDH2-R172K (ab264052, Abcam, 1:1000), anti-RARα (gifted by Prof. Hugues de Thé), anti-RXRα (3085, CST), anti-Actinin (3134, CST, 1:10000), anti-VDR (12550, CST).

2.7. Analysis of myeloid differentiation
2.7.1. Flow cytometry

We performed FACS analysis of CD11b, CD14 and CD15 expression using the following anti-human monoclonal antibodies: CD11b-PE (Beckman Coulter, IM2581U), CD14-AF700 (BD, 557923), CD15-APC (BD, 551376), CD45-BV510 (BD, 563204) or CD45-AFCH7 (BD, 641417). Apoptotic cells were stained with Annexin-V-BV421 (BD, 563973) or Annexin-FITC (BD, 556420) according to the manufacturer’s protocol (BD).

2.7.2. Morphological characterization

For the morphological analysis of myeloid cell differentiation, we prepared cytopsins by centrifugation of 200µL of PBS1X containing 50000 cells (800rpm, 10min for primary cells and 500rpm, 4min for clones from HL60 cell line) using SuperfrostPlus positively charged glass slides. Then, we stained cytopspun slides at room temperature with May-Grünwald Giemsa. AxiCam camera (Carl Zeiss) was used for image acquisition.

2.8. Data exploration mining

List of differentially expressed mRNA (FC> ±1.5 and p-value <0.05) for each cohort (Table S1) were uploaded in the Genome Analyzer bioinformatics tool (Genomatix) for pharmacological substances gene signatures enrichment analyses based on the Genomatix literature mining (Table S2-3). GSEA analyses was performed using GSEA version 3.0 (Broad Institute). The “Late VD/VDR activation pathway” gene signature used in this study were obtained from Warwick et al.[14]. Transcriptome of HL60 IDH1WT, IDH1WT+2HG, IDH1R132H, IDH1R132H+ATRA were generated by our team[4]. Publicly accessible transcriptomic databases used in this study: GSE144684, TCGA.

2.9. Single cell RNA sequencing

Viable primary human AML cells from mice bone marrow of PDX were isolated using cell sorter cytometer. Viable AML cells were then diluted to 1500 cells/µl in PBS1X 0.05. To generate single-cell libraries the Chromium Single Cell 3’ Reagent Kits (v3 Chemistry, 10x Genomics) was used. About 10,000 cells were added to the channel and sample was run according to manufacturer’s protocol with a targeted cell recovery estimate of
12,000 cells in total. After reverse transcription, single-cell droplets were broken and the single-strand cDNAs were isolated and cleaned. cDNAs were then amplified 12 thermal cycles. The sequencing-ready library was cleaned up prior to sequencing on a NextSeq550 instrument (Illumina). The output .bcl files were demultiplexed to FASTQ format by using bcl2fastq. FASTQ reads were aligned and converted into gene expression matrices with CellRanger-3.0.2. Expression matrices were loaded in an R session with the Seurat 4.0 package [15], which facilitated the normalization and variance stabilization of the data using the sctransform function [16]. For quality control purposes, only viable cells with a mitochondrial RNA expression ratio of 0.2 or below were maintained. Further filtering parameters include the exclusion of cells with unique molecular identifiers (UMIs) less than 500, number of genes less than 250, and log10(genes per UMI) less than 0.80. To visualize and infer common gene expression with minimal batch effect, samples were integrated by status of treatment (diagnosis, MRD, and relapse). Cell types were categorized with the “Hay Bone Marrow” signature collection from the Human Cell Atlas. After applying the AddModuleScore function from Seurat to derive signature expression scores per each cell per signature, the expression values were normalized to z-scores. The greatest z-score pertained to the highest-likelihood cell type by which it was then categorized. Z-scores for cellular function signatures were ascertained using the same method and visualized using the FeaturePlot function. The plotting of the correlation among signature expressions was facilitated with the FeatureScatter Seurat function, and statistical analyses to validate these correlations were provided by the base-R function cor.test.

2.10. Metabolite quantification

2.10.1. Extraction of metabolites

Metabolite extraction was performed from frozen cell pellets (-80°C). Briefly, pellets were mixed with 150 µL of water, and 5 µL of an external standard mixture containing 20µg/mL of 

$^{13}$C$_5$-2-HG, $^{13}$C$_5$-α-KG were added. Samples were scrapped, extracted with 350 µL of ice-cold methanol and vortexed. Next, samples were centrifuged at 20,000g for 15 min at 4°C, and dried under a nitrogen stream using a Turbovap (Biotage) at 30°C. Finally, samples were resuspended in 50 µL of mobile phase A (see below), vortexed and centrifuged at 4°C for 5min at 20,000g. The resulting supernatants were then transferred into vials.

2.10.2. Extraction of metabolites

Metabolites LC-HRMS quantification of 2-HG and α-KG was performed using an Ultimate 3000 chromatographic system coupled with an Exactive mass spectrometer (Thermo Fisher Scientific, Courtaboeuf, France) fitted with an electrospray source. Chromatographic separation was performed on a Hypercarb column (3µm, 2.1 x 100mm; Thermo Fisher Scientific, Massachusetts, United States) maintained at 30°C. Mobile phase A consisted of an aqueous buffer of 5 mM of ammonium formate at pH 2.7, while mobile phase B was made of acetonitrile with 0.1% formic acid. Chromatographic elution was achieved at a flow rate of 300 µL/min. After injection of 10 µL of sample, elution started with an isocratic step of 0.5 min at 95% A, followed by a linear gradient from 95 to 5% of phase A in 4.5 min. The chromatographic system was then rinsed for 1 min at 95% B, and the run ended with an equilibration step of 2 min. The column effluent was directly introduced into the electrospray source of the mass spectrometer, and analyses were performed in the negative ion mode. Source parameters were as follows: capillary voltage, -2.5kV; capillary temperature, 350°C; sheath gas and auxiliary gas pressures set at 35 and 10 arbitrary units, respectively. The detection was performed from m/z 50 to 850 at a resolution of 50,000 (full width at half-maximum at m/z 200) using an AGC target of 3E6 and a maximum injection time of 500ms. 2-HG and α-KG were detected as deprotonated [M-H]- species at m/z 147.0299 and 145.0142, respectively. 2-HG and α-KG, were quantified
by isotope dilution using the corresponding $^{13}$C-labeled homologues as internal standards.

2.11. Statistical Analysis

GraphPad Prism Software version 9 (La Jolla, CA, USA) was used for the statistical analysis. The results were expressed as a mean ± SEM. Statistical analyses were performed...
using two-tailed (non-directional) Student’s t test with Welch’s correction. The significance is represented by stars in which * is p < 0.05, ** is p < 0.01, *** is p < 0.005 and **** is p < 0.001. Combination indexes were calculated using Compusyn software.

3. Results

3.1. Vitamin D receptor-related gene signatures are enriched in transcriptomes of IDH mutant cells

To identify novel therapeutic agents for the treatment of AML harboring IDH mutation, we first performed transcriptomic analyses of two published independent cohorts of AML patients (GSE14468[17], TCGA) and HL60 AML cell line expressing either IDH1R132H or IDH1WT (Supplemental Table S1)[4]. We identified four pharmacological substance-associated gene signatures enriched in both IDH1MUT AML patients, HL60 IDH1R132H and HL60 IDH1WT treated with 2-HG including ATRA and VD (Figure 1A and supplemental Table S2). Both ATRA and VD are differentiating agents known to induce granulocytic and monocytic differentiation in AML, respectively [18]. Among the enriched genes in IDH1MUT and IDH1WT+2HG AML cells associated with these two small molecules, we identified common genes related to monocytic and granulocytic differentiation (RXRa, ITGAM or CD11b, ITGAX or CD11c, CEBPa) (Figure 1B and supplemental Table S3). Using Gene Set Enrichment Analysis (GSEA), we further confirmed the enhanced gene expression of VDR pathway in IDH1R132H HL60 cells compared to wild-type isoforms (Figure 1C and supplemental figure S1A). Particularly, the VDR-specific gene signature of late VR response (generated from Warwick et al 2021 [14], supplemental Table S4) is further enriched in transcriptome of IDH1WT AML cells treated with exogenous 2-HG and in patients harboring IDH mutations (Figure 1C, and supplemental Figure S1B). Using scRNA-seq of human IDH1R132H AML blasts from a patient-derived xenografted (PDX) model in NSG mice, we confirmed this phenotype in vivo. Of note, the Late VD/VDR activation pathway gene signature was strongly correlated to gene signatures of IDH1MUT patients generated from two independent public cohorts in the PDX IDH1 R132H (Figure 1D). This correlation is independent from the cell types (supplemental Figure S1C). Finally, transcriptomes of relapsing patients post-IDH1MUT inhibitors (NCT01915498, NCT02074839; GSE153349[19]) are also enriched in VDR gene signatures compared to those from patients before receiving IDH inhibitors therapies (Figure 1E).

Taken together, transcriptomic analysis of cell lines, primary patient cells and PDX with IDH mutations showed that VD/VDR pathway is activated in this subgroup of AML patients. Furthermore, this VD/VDR gene signature is enhanced in patients who are relapsing from IDH inhibitor treatments, suggesting that Vitamin D-based therapies could represent a promising alternative for these patients.

3.2. IDH mutations activate CEBPa-VDR-RXR axis through 2HG production

Next, we analyzed the expression of VDR in HL60 expressing IDH1WT or IDH1R132H. Consistently, we observed an increased expression in VDR, RARα (ATRA receptor/transcription factor) and RXRα (VDR and RARα co-receptor/transcription factor) in IDH1R132H (Figure 2A,B). Then, we asked if this phenotype was the same upon IDH2 mutation. To assess this question, we generated HL60 cell line expressing IDH2WT or IDH2R172K (supplemental Figure S2A,B). As for IDH1 mutation [4], IDH2 mutation leads to an increase in CEBPa protein expression (supplemental Figure S2A). Importantly, both mutations enhanced VDR, RXRα and RARα protein expressions compared to the WT enzymes (Figure 2B). This increase in VDR, RARα and RXRα was also induced when IDH1WT HL60 cells were treated with 2-HG (Figure 2C,D).
Figure 2. IDH mutations activate CEBPα-VDR-RXRα axis through 2HG production. (A) RT-qPCR showing gene expressions of VDR and RARA in HL60 IDH1WT (clone 2: ○; clone 7: □) versus HL60 IDH1R132H (clone 5: ◇; clone 11: △). (B) Western blot showing protein levels of VDR, RXRα and RARα in HL60 IDH1WT (clones 2 and 7) versus HL60 IDH1R132H (clones 5, 11 and 3) (left) and in HL60 IDH2WT versus HL60 IDH2R172K (right). (C) RT-qPCR showing gene expressions of VDR and RARA in 2HG-treated HL60 (200 µM, 1 week). (D) Western blot showing protein levels of VDR RXRα and RARα in 2HG-treated HL60 IDH1WT (clones 2 and 7) (200 µM, 1 week). (E) RT-qPCR showing gene expression of and VDR and RARA in HL60 IDH1WT (clone 2: ○, clone 7: □) and IDH1R132H (clone 5: ◇, clone 11: △) after CEBPα-KD by shRNA (F) Western blot showing protein levels of CEBPα, VDR, RARα and VDR in HL60 IDH1WT (clones 2 and 7) and IDH1R132H (clone 5 and 11) after CEBPα-KD by shRNA. (G) Western blot showing protein levels of VDR in 2HG-treated HL60 IDH1WT (clone 2 and 7) with CEBPα-KD by shRNA (200µM, 1 week).

We have previously shown that 2-HG production in IDH1MUT AML cells leads to CEBPα activation through a histone methylation-dependent mechanism [4]. Moreover, CEBPα modulates vitamin D signaling in hematopoietic, AML and breast cancer cells [14], [20], [21]. Furthermore, recent studies have demonstrated that demethylating agents induce VDR expression in AML and in high CEBPα breast cancer cells [14], [22]–[24]. Thus, we determined whether VDR and RXRα might be downstream of CEBPα. Knockdown of CEBPα led to a decrease in VDR expression and RXRα expression in both IDH1WT and IDH1R132H cells, although the effect on VDR protein expression was markedly enhanced in IDH1MUT cells and RXRα decrease following CEBPα silencing was only observed in IDH1MUT cells compared to IDH1WT cells (Figure 2E,F and supplementary Figure S3A). Notably, 2-HG failed to increase VDR expression in IDH1WT cells invalidated for CEBPα, confirming the crucial role of the 2-HG/CEBPα/VDR axis in IDH1MUT AML (Figure 2G). Next, we investigated if other transcription factors known to regulate myeloid/monocytic maturation such as CEBPB might be enhanced by IDH mutation. Importantly, CEBPα and CEBPB are also direct targets of VDR-dependent transcriptional regulation [21]. However, CEBPβ expression was not modified either by IDH1 mutation or 2-HG-treatment (Supplementary Figure S3A,B,C), suggesting that VDR pathway enhancement is regulated through CEBPα rather than CEBPβ in IDH1MUT AML. Accordingly, CEBPβ invalidation by shRNA failed to decrease VDR and RXR protein expression (Supplemental Figure S3A). Overall, these results showed that 2-HG production and IDH mutations activate CEBPα, leading to an increase in VDR/RXRα pathway.

3.3. Targeting vitamin D receptor pathway enhances anti-AML effect with ATRA in a CEBPα-dependent manner especially in IDH1MUT cells.

Lastly, transcriptomes of IDH1R132H HL60 treated with ATRA were also enriched in genes related to VD (Supplementary Figure S4A and supplemental Table S5), suggesting a putative synergistic effect of these two differentiating agents in this cellular context. To further explore the therapeutic benefit of these agents for AML patients with IDH mutation, we treated primary IDH1MUT AML cells with ATRA or VD alone or in combination. We analyzed mono-granulocytic marker expression by flow cytometry in AML cell lines (CD11b) and primary patient cells (CD11b, CD15, CD14) with IDH1WT or IDH1MUT. We first observed that the combination therapy induced a higher differentiation potentiation as shown with an increased CD11b expression in IDH1R132H and IDH2R172K HL60 cells compared to IDH1WT HL60 cells (Figure 3A and supplemental Figure S4B). Combination index analysis showed that VD and ATRA acted synergistically to induce CD11b differentiation markers expression in IDH1R132H HL60 cells (Figure 3B,C). Then, morphological analysis confirmed the enhancement of myeloid differentiation characterized by an increase in number of monocyte-like cells in IDH1R132H AML cells upon doublet therapy (Figure 3D).
Figure 3. Targeting vitamin D receptor pathway enhances anti-AML effect with ATRA in a CEBPα-dependent manner. (A) HeatMap of CD11b expression (MFI, ratio to untreated) measured by flow cytometry in HL60 IDH1<sup>WT</sup> versus HL60 IDH1<sup>R132H</sup> and HL60 IDH<sup>2WT</sup> versus HL60 IDH2<sup>R132K</sup> treated for 3 days with ATRA (1µM) and VD (100nM) alone or in combination. (B) Synergy Map of CD11b expression (MFI, ratio to untreated) measured by flow cytometry in HL60 IDH1<sup>R132H</sup> (left) and in HL60 IDH<sup>R132H</sup>-5<sup>11</sup> (right) treated for 3 days with 0.25-4µM of ATRA and 25-400nM of VD alone or in combination. (C) Combination index of differentiation (CD11b expression induction) in HL60 IDH1<sup>R132H</sup>-5<sup>11</sup> and in HL60 IDH1<sup>R132H</sup>-11 treated for 3 days with ATRA and VD in combination with constant ratio of each drug. (D) Morphological characterization by MGG staining of HL60 IDH1<sup>WT</sup>-2 versus HL60 IDH1<sup>R132H</sup>-11 treated for 5 days with ATRA 1µM and VD 100nM alone or in combination. (E) HeatMap of CD11b expression and CD15 expression (MFI, ratio to untreated) measured by flow cytometry in 2HG-treated (100(U937)-200µM for 1wk) HL60 IDH1<sup>WT</sup>-2, HL60 IDH1<sup>WT</sup>-4, HL60 IDH1<sup>WT</sup>-7, HL60, U937, THP1 and KG1a treated for 3 days with ATRA (0.1µM for U937, 1µM for others) or VD (100nM) alone or in combination. (F) CD11b expression (MFI) measured by flow cytometry in HL60 IDH1<sup>WT</sup> (clone 2: ○, clone 7: □) shCTL vs. shCEBPα (left panel) and in HL60 IDH1<sup>R132H</sup> (clone 5: ○, clone 11: △) shCTL vs. shCEBPα (right panel) treated for 3 days with ATRA (1 µM) and VD (100nM) alone or in combination. (G) CD11b expression (MFI) measured by flow cytometry in 2HG-treated HL60 IDH1<sup>WT</sup> (clone 2: ○, clone 7: □) shCTL vs. shCEBPα (2HG 200µM for 3 days) treated for 3 days with ATRA (1 µM) and VD (100 nM) alone or in combination.

Consistent with the enhanced activation of the VDR and ATRA pathways in HL60 IDH1<sup>WT</sup>-2HG (Figure 2C,D), 2-HG pre-treatment of IDH1<sup>WT</sup> and four parental AML cell lines elicited a higher sensitivity to differentiation induction by ATRA/VD combination (Figure 3E and supplemental Figure S4C,D). Furthermore, CEBPα knockdown reduced significantly the response to this combination ATRA/VD in IDH1<sup>R132H</sup> cells while no difference was observed in IDH1<sup>WT</sup> cells (Figure 3F). Interestingly, CEBPβ knockdown also reduced the efficiency of the combination on the differentiation of IDH1<sup>R132H</sup> AML cells but at a significant lower extent (supplemental Figure S4E). Finally, we showed that, in the absence of CEBPα, 2-HG no longer enhanced sensitivity to ATRA/VD combination-induced differentiation of IDH1<sup>WT</sup> cells (Figure 3G).

Despite phenotypical heterogeneity between patients, we also observed a marked enhancement of differentiation induction when ATRA was used in combination with VD in 13 out of 17 primary IDH<sup>MUT</sup> AML cells compared to 10 primary IDH<sup>WT</sup> AML cells (Figure 4A; supplemental Figure S5; patient characteristics are summarized in Supplementary Table S6). Morphological characterization of AML blasts from Ps6 using MGG staining confirmed the analysis of myeloid differentiation markers with an increase number of granulocytic/monocytic-like cells upon ATRA/VD (Figure 4B). Then, we classified IDH<sup>MUT</sup> patients according to the efficiency of ATRA and VD co-treatments to enhance differentiation when they were associated. We clustered the patients in 4 following groups: CD11b/CD15 responders (n=7), CD11b responders (n=4), CD15 responders (n=2) and the non-responders (n=4) (Figure 4A). Next, we compared these groups with the clinical characteristics of these IDH<sup>MUT</sup> patients. We did not observe any correlation between the differentiation response and other mutations or FAB status. However, we noted that, among responders to ATRA/VD, we had a large proportion of patients with an adverse or intermediate risks according to ELN2017 stratification [25] (Figure 4C). This result suggests that ATRA/VD combination therapy represents a promising therapeutic approach for IDH<sup>MUT</sup> AML patients with a poor prognosis.
Figure 4. Targeting vitamin D receptor pathway enhances anti-AML effect with ATRA in IDH\textsuperscript{MUT} AML patients. (A) CD11b and CD15 expressions (mean fluorescence intensity, MFI) measured by flow cytometry in IDH\textsuperscript{MUT} versus IDH\textsuperscript{WT} patients treated with ATRA and VD alone or in combination (left panel). Results for each patient are represented individually (right panel). (B) Morphological characterization by MGG staining of Ps6 treated for 14 days with ATRA 0.1\textmu M and VD 50-100nM alone or in combination. (C) Clinical characteristics of IDH\textsuperscript{MUT} patients in function of their response to ATRA/VD-doublet therapy

4. Discussion

The past few years have witnessed major advances in the therapeutic landscape of AML treatment [7]. Translational research has led to a better understanding of the impact of multiple recurrent mutations, leading to the successful development and approval of targeted therapies in several subgroups of patients, including the ones harboring IDH mutations [26]. While small-molecule IDH\textsuperscript{MUT} inhibitors allow promising partial or complete responses rates in AML patients, about 60% of them do not respond to the therapy or relapse [27]–[30]. Very importantly, 2-HG level is reduced in nearly all patients treated with IDH\textsuperscript{MUT} inhibitors, including non-responders, revealing that 2-HG reduction is not sufficient to predict clinical response to these inhibitors [28], [31]. This observation may also indicate that the ability of primary patient cells to elicit alternative resistance mechanisms will dictate the clinical benefit of the IDH\textsuperscript{MUT} inhibitors. Some of these mechanisms notably induce a 2-HG rebound through second-site mutation or isoform-switching [9], [32]. Exploiting the 2-HG specific effects in IDH\textsuperscript{MUT} AML cells could therefore be an alternative therapeutic approach for patients who are non-responding or relapsing from IDH\textsuperscript{MUT} inhibitors treatments. In this study, we demonstrated that 2-HG mediates the activation of VDR pathway through CEBP\textalpha in IDH\textsuperscript{MUT} AML, sensitizing IDH\textsuperscript{MUT} AML cells to the differentiation-based ATRA/VD doublet therapy. In this context, we proposed the association of VD with ATRA -both already used in clinic- as a promising approach to induce terminal differentiation of leukemic cells in IDH\textsuperscript{MUT} AML patients.

Numerous studies have investigated the potential benefit of VD-induced differentiation through interaction with VDR in different cancers including AML, without encouraging results [33]. Indeed, regulation of VDR is complex and highly dependent on the cellular context. AML cell line KG1 has been shown to be resistant to VD-induced monocytic differentiation but sensitive to ATRA-induced granulocytic differentiation, while HL60 displayed inverse responses to these two agents [34]. ATRA treatment enhances VDR expression in KG1 cells, resulting in increased differentiation using the ATRA+VD combination. As opposed, ATRA treatment reduces VDR gene expression in HL60 cells, leading to no better response to the ATRA+VD combination [34]. In our study, we showed that IDH\textsuperscript{MUT} through production of 2-HG increases VDR expression, which could therefore explain why VD addition to ATRA induced even more differentiation in all the different cell lines tested, including notably HL60. Interestingly, VD induced CEBP\textalpha in breast cancer MCF-7 cells and CEBP\textalpha knockdown suppressed VD anti-proliferative effects. As opposed, in breast cancer MDA-MB-231 cells non-responsive to VD treatment, CEBP\textalpha was not detected and CEBP\textalpha over-expression restored VD sensitivity [21]. Additional studies demonstrated that VD rather induced CEBP\beta in kidney and osteoblasts cells [35] and in HL60 AML cell line [36], for which CEBP\beta silencing inhibited VD-induced differentiation [37]. Of note, exogenous 2-HG can inhibit proliferation and viability in some leukemia cell lines [38]. In these sensitive IDH\textsuperscript{WT} AML cells, 2-HG targets the fat mass and obesity-associated protein (FTO), that leads to MYC/CEBP\textalpha inhibition. However, most IDH\textsuperscript{MUT} AML cells upregulate MYC which counteracts the effect of 2-HG on proliferation, CEBP\textalpha expression and metabolic consequences [38], [39]. Indeed, for AML cells harboring IDH mutation, myeloid lineage-TF CEBP\textalpha is one of key downstream pathways that are critical for IDH mutant AML biology [4], [13]. In these cells, the activation
of CEBPα induces the expression of both RXRα and VDR, that primes to VD+ATRA combination therapy. In particular, 2-HG-induced epigenetic changes (e.g., histone methylation) might affect the chromatin accessibility status and structure, favoring VDR DNA binding with pioneer factors such as PU.1 or and CEBPα [20], [40]. Noteworthy, Warwick et al (2021) showed that the regulatory network of VD response is associated with 47 transcription factors and that its transcriptional hierarchy is specifically and epigenetically dependent on VDR and CEBPα in monocytic AML THP1 cells [14]. CEBPα silencing downregulates and inactivates more than half of VD target genes [14]. Therefore, further similar molecular studies should be performed to identify new transcriptional partners and VD targets specific to IDH mutant context. Markedly, Glass at al. showed a differential CpG methylation in RXR promoter specifically in IDH2MUT AML cells [41]. Overall, IDH mutations seems to profoundly disrupt the epigenetic landscape of AML cells leading to the acquisition of a specific transcription factor regulation network responsible of their sensitivity to differentiating agents. Here, we demonstrated that CEBPα is crucial to ATRA and VD-induced differentiation response as knockdown of CEBPα, and CEBβ to significant lower extent, decreased IDH M differentiation and almost completely abrogated differentiation in IDH WT treated with exogenous 2-HG. Of note, high serum 2-HG and low serum VD levels are associated with shorter overall survival [43]–[45], they therefore might represent potential biomarkers to predict response to ATRA/VD doublet therapy for IDH M AML patients. Although randomized clinical trials are still lacking, epidemiological and preclinical data suggest that VD-based combinations could be a promising therapeutic strategy by activating VD signaling pathway. In this context, better designs of combinations would circumvent potential limitations of VD alone such as high risk of hypercalcemia.

Finally, hypomethylating agents such as 5-azacytidine (AZA) or decitabine induce an increase in expression of VDR and its target genes in breast cancer cells (Marik et al, [22]) and AML cells [23]. In vivo treatment of leukemic mice with VDR agonist INEC, AZA alone, and in combination led to no difference in mice survival for AZA alone and to a significant but similar increase in mice survival for INEC alone and INEC+AZA [23]. As AML patients with IDH mutations seem to have better response to hypomethylating agents [42], combination of VD and ATRA with hypomethylating agents could also represent interesting combinations. A current recruiting phase 2 clinical trial combining INEC and decitabine in AML patients unfit for standard chemotherapy (NCT02802267) could help determine if patients with IDH mutations are more likely to respond. If so, further studies measuring the expression of VDR and VD target genes, as well as differentiation and impact on the stem cell compartment specifically in IDH M AML cells following these combinations would be of particular interest and help determine potential biomarkers of response.

5. Conclusions

In conclusion, these data demonstrate that the VDR pathway is increased in IDH M subgroup, especially at relapse post-IDHm inhibitor. Moreover, VD could enhance ATRA effect in AML patients with IDH mutation through a 2-HG/CEBPα/VDR axis. Thus, our study describes an alternative approach with approved differentiation drugs ATRA and VD with a highly efficient and safety profile, already in the clinics. This strategy could be specifically used for non-eligible AML patients to IDH mutant inhibitor therapies or AML patients developing resistance to newly FDA-approved IDH mutant-specific inhibitors. This is timely and clinically relevant and forms the basis of novel doublet and triplet therapies in IDH-mutated AML patients.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Figure S1: Vitamin D receptor-related gene signatures are enriched in transcriptomes of IDH mutant cells, Figure S2: Generation of HL60 expressing IDH2MUT or IDH2T27K, Figure S3: IDH mutations activate CEBPα-VDR-RXRα axis through 2HG production, Figure S4: Targeting vitamin D receptor pathway enhances anti-AML effect with ATRA in a CEBPα-dependent manner, Figure S5: Targeting vitamin
D receptor pathway enhances anti-AML effect with ATRA in IDH\textsuperscript{MUT} AML patients, Table S1: Gene signatures related to Figure 1A, Table S2: List of pharmacological substances from VENN diagram related to Figure 1A, Table S3: ATRA- and VD-responsive genes enriched in IDH\textsuperscript{MUT}-AML and HL60WT +2HG, Table S4: Gene signatures used for GSEA analysis, Table S5: Pharmacological substances (Genomatix) enriched in gene signature of Atra-treated HL60 IDH1R132H (HR132H) related to Figure 1M, Table S6: Clinical characteristics of AML patients used in this study and related to Figure 4.


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