Leukemia cells remodel marrow adipocytes via TRPV4-dependent lipolysis

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Introduction

The development of acute myeloid leukemia (AML) is closely related to the bone marrow (BM) microenvironment. As a critical component of the BM microenvironment, BM adipocytes provide energy for both the infinite proliferation of leukemia cells and the normal growth of hematopoietic stem cells. Leukemia cells proliferate to an overwhelming number in a limited marrow cavity, likely because these cells are more efficient in capturing energy for growth. Accordingly, BM adipocytes are remodeled in response to leukemia cells, generating a pro-tumoral microenvironment. However, the mechanism whereby leukemia cell growth induces BM adipocyte remodeling is still unclear.

Induced BM adipocyte remodeling involves several specific processes, including lipolysis, dedifferentiation and lipid accumulation. Consequently, the remodeled adipocytes show morphological and functional changes. Breast cancer cells reportedly secrete soluble factor Wnt5a which reduces the number and size of adipocytes surrounding the malignant cells and thus contributes to disease deven-
As the breast cancer progresses, adipocytes de-differentiate to fibroblast-like cells. In mouse models of bone metastasis following prostate cancer, Herroon et al. showed that remodeled BM adipocytes support tumor growth by fatty acid-binding protein 4 (FABP4) trans- portation of fatty acids. These studies identified a func- tional role of remodeled adipocytes in supporting solid tumor metabolism. Thus, the adipocytes remodeled by cancer cells are also known as cancer-associated adipocytes. In the context of leukemia, there is a grow- ing consensus that reduction in BM adipocyte number, once believed to be merely due to mechanical squeezing by the rapid proliferation of leukemia cells in the limited BM cavity, is also actively regulated by leukemia cells. Indeed, we previously reported that growth differentia- tion factor 15 (GDF15) derived from leukemia cells regul- ates BM adipocyte remodeling by enhancing lipolysis. However, how extracellular GDF15 induces lipolysis within BM adipocytes remains elusive.

It has been reported that GDF15 enhances intracellular Ca2+ by increasing calcium voltage-gated channel subunit alpha1 C (Cav1.3) expression in rat cerebellar granule neurons, which induces the expression of genes essential for synaptic plasticity. As an important cellular signal for lipid metabolism, intracellular Ca2+ is involved in lipid synthesis and lipolysis in adipocytes. When the calcium channels in the adipocytes are activated or upregulated, accumulation of lipids is enhanced through increased [Ca2+]i. Conversely, when calcium channels are inhib- ited or downregulated, decreased calcium influx may accele- rate fat breakdown. Thus, we hypothesized that calcium channels are involved in GDF15-induced BM adipocyte remodeling.

In this study, we examined a possible role of transient receptor potential vanilloid 4 (TRPV4) calcium channels in GDF15-driven remodeling of BM adipocytes. We unravel a novel function of transforming growth factor-β type II receptor (TGFβRII) that, in responding to GDF15 in BM adipocytes, activates the phosphatidylinositol 3- kinase (PI3K)/AKT transduction pathway, which in turn reduces the transcript factor Forkhead box C1 (FOXC1) level and subsequently downregulates TRPV4. We also provide evidence that inhibition of BM adipocyte remodeling increases survival in the AML mouse model, implying a novel therapeutic target for AML.

Methods

Patients’ samples

BM aspirates were collected from 16 patients diagnosed as having lymphoma without BM invasion, using procedures approved by the Ethics Committee of Shanghai Jiao Tong University Affiliated Hospital. Mesenchymal stem cells were derived from the BM of lymphoma patients without BM inva- sion, because marrow mesenchymal stem cells in this type of patients can be considered normal. The adipogenic induction of mesenchymal stem cells is described in the Online Supplementary Methods.

Chromatin immunoprecipitation-quantitative polymerase chain reaction

Adipocytes were collected from different groups and crosslinked with 1% formaldehyde for 10 min at 37°C. Cross- linking was blocked, then the cells were washed and lysed in sodium dodecylsulfate lysis buffer (50 mM HEPE NaOH 7.5, 500 mM NaCl, 1 mM EDTA, 0.1% Na-deoxycholate, 1% Triton X100). The lysates were sonicated to shear DNA to a length between 200 and 500 base pairs with 10-second pulses using sonication. The antibody against FOXC1 (5 μL, Abcam5079, USA) was then added to the supernatant, incubated overnight at 4°C with rotation and incubated with 100 μL Salmon Sperm DNA/Protein A agarose beads for 2 h at 4°C. The immunoprecipitated complex was then washed and eluted. The histone DNA crosslinks were reversed and DNA was purified for real-time poly- merase chain reaction (PCR). Quantitative real-time PCR (RT-qPCR) was performed on bound and input DNA with the following primers for TRPV4: forward: 5-CTTTGCACTGGGAGCA-GAGT-3, reverse: 5-ATTAGCGTGGGTTACAGCA-3.

Cell cultures and reagents

The cell cultures and reagents, as well as the co-culture assays are described in detail in the Online Supplementary Methods.

Animal experiments

All animal experiments were performed according to proce- dure approved by the Ethics Committee of Shanghai Jiao Tong University Affiliated Hospital. Five-week old C57BL/6 mice were fed with 60% high-fat diet (Research Diets, Inc. New Brunswick, NJ, USA) for 3 months to create an obese mouse model. Mice injected with FBL-3 cells (5x10^5) and mice injected with both FBL-3 cells (5x10^5) and 4α-phorbol 12,13-didecanoate (4αPDD) (200 μg/Kg according to the instructions for reagents) were used as experimental groups. The untreated obese mice were used as a control group. The volume of all solutions inject- ed was 200 μL. Mice were sacrificed and femora were removed after 3 weeks of treatment. Femora were fixed for 24 h with 4% paraformaldehyde and were decalcified for 2 days. BM sections from the mice were dewaxed by conventional methods and incubated with anti-perilipin1 monoclonal antibody (1:50, CST, USA) at 4°C overnight.

Other experimental details

Full descriptions of free fatty acid detection, lentiviral knock- down, RNA sequencing, western blot analysis and enzyme- linked immunosorbent assays, RT-qPCR (primers shown in Table 1), the cell counting and apoptosis assays, oil red O staining, immunofluorescence studies, and adipocyte measurements are provided in the Online Supplementary Methods.

Statistical analysis

All statistical tests were performed with GraphPad Primer5. The data are presented as the mean ± standard deviation. A Student t-test was used for comparisons between two groups. A P value of less than 0.05 was considered statistically significant.

Results

Downregulated TRPV4 contributes to increased bone marrow adipocyte lipolysis

As an important channel for calcium ions, TRPV plays a critical role in the energy balance of adipocytes. RT-qPCR analysis showed that TRPV4 mRNA in BM adipocytes had the highest expression among TRPV family members (Figure 1A). Moreover, western blot analysis showed that BM adipocytes expressed TRPV4 protein highly (Figure 1B). To investigate whether TRPV4 plays an important role in BM adipocytes, we used TRPV4 inhibitor (RN1734) and agonist (4αPDD) to verify the
Figure 1. Downregulated TRPV4 contributes to increased bone marrow adipocyte lipolysis. (A) Reverse transcriptase quantitative polymerase chain reaction (RT-qPCR) verification of the expression of transient receptor potential vanilloid (TRPV) channel genes in bone marrow (BM) adipocytes. (B) Western blot analysis of TRPV4 protein in BM adipocytes from three patients. (C) BM adipocytes treated with dimethylsulfoxide (Ctr), RN1734 (5 μM) or 4α-phorbol 12,13-didecanoate (4αPDD, 0.25 μg/mL) for 4 days. Adipocytes were stained by oil red O (ORO). All images were at a magnification of 200×. (D) The number and average area of BM adipocytes from the indicated groups were measured using Image-Pro-Plus 5.1. (E) The content of lipid droplets in BM adipocytes from the indicated groups was detected by optical density values after ORO staining. (F) RT-qPCR was used to analyze adipose triglyceride lipase (ATGL) and hormone-sensitive triglyceride lipase (HSL) mRNA in BM adipocytes from the indicated groups. (G) The content of free fatty acids (FFA) in the supernatant of BM adipocytes treated with dimethylsulfoxide (Ctr), RN1734 or 4αPDD was detected using the colorimetric method. (H) BM adipocytes were infected with TRPV4-targeted shRNA (shTRPV4) lentivirus for 6 days. Adipocytes were stained with Alexa Fluor 493/503-conjugated BODIPY. 4,6-diamidino-2-phenylindole (DAPI) stained blue and lipid droplets showed green fluorescence. The scale bar represents 50 μm. (I) The number and area of BM adipocytes infected with shTRPV4 lentivirus, quantitatively analyzed by Image-Pro-Plus 5.1. (J) The mRNA level of HSL and ATGL in BM adipocytes infected with shTRPV4 lentivirus on the fourth day, detected by RT-qPCR. β-actin protein was used as an internal control for the western blot analysis. Three independent experiments were performed. ***P<0.001, **P<0.01, *P<0.05.
function of TRPV4 in BM adipocytes, respectively. 4αPDD is the first synthetic TRPV4 agonist and is a non-protein kinase C activated phorbol ester. Online Supplementary Figure S1A shows the half maximal inhibitory concentration (IC50) of the effects of RN1734 on BM adipocytes. Considering excessive Ca2+ influx could cause some toxicity to adipocytes, we aimed to find a concentration that has minimum cellular toxicity and promotes the Ca2+ influx needed for our experiments. As Online Supplementary Figure S1B shows, 4αPDD at a concentration of 0.25 μg/mL resulted in an acceptable level of toxicity of adipocytes, while allowing Ca2+ influx to reach the level required for the experiment. Oil red O staining and quantitative analysis showed that RN1734 reduced the number and area of BM adipocytes, whereas 4αPDD did not induce a similar change (Figure 1C, D), suggesting that the inhibition of TRPV4 contributes to reducing BM adipocyte number and size. Furthermore, optical density value measurements showed that lipid droplets in BM adipocytes treated with RN1734 decreased significantly (Figure 1E).

In order to determine whether the phenomenon is related to lipolysis, we determined the rate-limiting enzymes (adipose triglyceride lipase, ATGL and hormone sensitive lipase, HSL) of lipolysis. ATGL catalyzes the first step of lipolysis and converts triglyceride to diacylglycerol and free fatty acids. HSL is a hydrolase of glyc erides and cholesterol esters. Along with TRPV4 channel inspiration, BM adipocytes subsequently exhibited increased expression of ATGL and HSL, which resulted in increased free fatty acids in the supernatant (Figure 1F, G). Furthermore, it was found that RN1734 could significantly inhibit Ca2+ influx in BM adipocytes, while 4αPDD promoted Ca2+ influx in BM adipocytes (Online Supplementary Figure S1C). However, 4αPDD activates calcium channels in BM adipocytes by promoting Ca2+ influx, while the expression of TRPV4 could not increase (Online Supplementary Figure S2A).

To further confirm that TRPV4 regulates lipolysis of BM adipocytes, we used shTRPV4 lentivirus to knock down TRPV4 (Online Supplementary Figure S2B, C). As shown in Figure 1H, I and Online Supplementary Figure S2D, quantitative analysis showed that the number (control vs. shTRPV4, 528.1±46.4/mm² vs. 298.9±48.3/mm², P<0.05) and area (control vs. shTRPV4, 798.7±57.5 μm² vs. 454.7±54.0 μm², P<0.01) of BM adipocytes decreased in TRPV4 knockdown samples. ATGL and HSL mRNA levels were also increased in TRPV4 knockdown adipocytes (Figure 1J). These data indicate a critical role for TRPV4 in the regulation of lipolysis in BM adipocytes.

### TRPV4 mediates GDF15-induced bone marrow adipocyte remodeling

Increased lipolysis can result in a decrease in the number and area of BM adipocytes. Therefore, lipolysis is also a form of adipocyte remodeling. Our previous studies found that GDF15 secreted by leukemia cells promoted BM adipocyte lipolysis, decreasing the number and area of BM adipocytes. As shown by western blot analysis, TRPV4 expression was inhibited in BM adipocytes when co-cultured with leukemia cell lines (THP-1, K562, HL-60), whereas anti-GDF15 neutralizing antibodies partly reversed the effect (Figure 2A). Given the above results, we added recombinant human GDF15 (rhGDF15) to BM adipocytes to clarify this effect. It was shown that the inhibitory effect on TRPV4 was enhanced with the increase of rhGDF15 concentration and treatment duration (Figure 2B, C). Moreover, rhGDF15 could significantly inhibit TRPV4 mRNA expression and increase pHSL protein expression on the fourth day (Figure 2C and Online Supplementary Figure S2E). However, the pHSL protein and the release of free fatty acids did not increase sig-

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**Table 1. Sequences of the primers used to detect gene expression by reverse transcriptase quantitative polymerase chain reaction.**

| Species | Name         | Forward                     | Reverse                      |
|---------|--------------|-----------------------------|------------------------------|
| human   | GAPDH        | CGAGCCGTCACAGGATTTGTCGTAT   | AGGTCTTCTCATGTGTTGAAAGAC     |
| human   | TRPV1        | CAGGCTTTATAGCTGCAGGAG       | TTTGACCTGGTAGTCTGAGGAAG      |
| human   | TRPV4        | CGTCCAACTGGGATGAGGTTACCCT   | CTTCCATCTCTGGTGACCTGT       |
| human   | TRPV5        | GGTGTTCTACCTAAGGCAGAG       | CTCAGCAAGCAGGAGAAGCT       |
| human   | TRPV6        | ACTGGTCCCCGTCCTCTGAGC       | GTGGTGAAGATAATGTCGCAAGC     |
| human   | ALK4         | GCCATGCGGAGGTTGATAGGG       | GCTGCAGGTCCGTAACTCACAG      |
| human   | ACVR2        | GCCACCTTATACCAACATCCTG      | GTGTGCGGTTGCTGAGTGG         |
| human   | TGFβRII      | GAGAGGACCCCTCATTGAG         | TGGCTACCGGACACGACGAGC       |
| human   | TGFβRIII     | CCAACACATCCCAACACA          | TTATAAGCCCTGAGAACAGC        |
| human   | GFRAL        | ATGCATCAAGAGGATGGA         | TGATAGAAACGACGTAGGCC        |
| human   | ATG1         | GGCGGTCGAGGAGCGAGAATG       | GCAGCTTCTGGAAGTGTTGAG       |
| human   | HSL          | CACATCAAACGCAACGACAGC       | CCAAGACGATAGCACTCCC         |
| human   | FOXC1        | TAAGCGCAATGACGCGC          | GCCGACGCTTCCCTACTCT         |
| human   | Cav3.1       | CTCCTCTCCTCATTCTACCTCTC    | TCTATCATTCCGGATCC           |
| human   | Cav3.2.1     | CCACAGGGTGCTCTGCTACTCA     | GGCTAGGAGGAGATAGGTCCTAGA    |
| human   | Cav3.2.2     | TCAGGAGGGACCTCCACAGG        | TGATACCGGAGGAGTGGG          |

GAPDH: glyceraldehyde 3-phosphate dehydrogenase; TRPV: transient receptor potential vanilloid; ALK4: activin A receptor B; ACVR2: activin receptor type 2; TGFβRII: transforming growth factor β type II receptor; FOXC1: forkhead box C1; Cav3.1: calcium voltage-gated channel subunit alpha1 G; Cav3.2: calcium voltage-gated channel subunit alpha1 H; Cav3.3: calcium voltage-gated channel subunit alpha1 I.
Figure 2. TRPV4 mediates GDF15-induced bone marrow adipocyte remodeling. (A) Bone marrow (BM) adipocytes were co-cultured with leukemia cell lines (THP-1, K562, HL-60) or leukemia cells and anti-GDF15 neutralizing antibody (200 ng/mL) for 4 days. The protein of TRPV4 was detected using western blot analysis. (B) The effect of different concentrations (100 ng, 200 ng, 500 ng) of recombinant human GDF15 (rhGDF15) on the expression of TRPV4 protein for 4 days was analyzed by western blot. (C) Reverse transcriptase quantitative polymerase chain reaction (RT-qPCR) was performed to analyze the expression of TRPV4 mRNA after the addition of 200 ng rhGDF15 in BM adipocytes for 2, 4, 6, 8 days. (D) BM adipocytes were treated with dimethylsulfoxide (Ctr), rhGDF15 (200 ng/mL) or rhGDF15 (200 ng/mL) and 4α-phorbol 12,13-didecanoate (4αPDD, 0.25 μg/mL) for 6 days. Adipocytes were stained by oil red O. All images were at a magnification of 200×. (E) The number and average area of adipocytes from the indicated groups were measured using Image-Pro-Plus 5.1. (F) The content of lipid droplets in the indicated groups was detected by optical density values. (G) RT-qPCR was used to analyze HSL and ATGL mRNA in adipocytes from the indicated groups on the fourth day. (H) The content of free fatty acids in the supernatant of BM adipocytes from each group was detected using a colorimetric method. β-actin protein was used as an internal control for the western blot analysis. Three independent experiments were performed. **P<0.01, *P<0.05.
significantly after BM adipocytes were treated with rhGDF15 for 1 h (Online Supplementary Figure S2F, G), suggesting that the role of GDF15 in promoting lipolysis may be different from the rapid action of β-adreneline. Oil red O staining and quantitative analysis showed that the number and area of adipocytes did not change significantly on the fourth day, but decreased significantly from the sixth day (Online Supplementary Figure S2H, I). These results indicate that TRPV4 regulates the remodeling of BM adipocytes.

To further explore the potential role of TRPV4 in GDF15-induced BM adipocyte remodeling, oil red O staining and quantitative analysis was conducted and showed that the number and area of BM adipocytes were decreased in BM adipocytes treated with rhGDF15, whereas 4αPDD partly reversed the effect of rhGDF15 (Figure 2D, E). Optical density value measurements of lipid droplets showed similar results (Figure 2F). Consequently, rhGDF15 could induce increased expression of lipolysis genes (ATGL and HSL) and increased release of free fatty acids from BM adipocytes, but activation of TRPV4 by 4αPDD partly reversed the effect of rhGDF15 (Figure 2G, H). Furthermore, rhGDF15 can inhibit Ca²⁺ influx in BM adipocytes (Online Supplementary Figure S3A). These findings strongly suggest that TRPV4 contributes to GDF15-induced remodeling of BM adipocytes. Although GDF15 has been reported to act on Cav1.3, Cav3.1, Cav3.2, Cav3.3, the expression of these channels in BM adipocytes is much lower than that of TRPV4 (Online Supplementary Figure S3B). Moreover, when BM adipocytes were co-cultured with leukemia cell lines (THP-1, K562, HL-60), the expression of TRPV4 changed significantly (Online Supplementary Figure S3C). These results further suggest that TRPV4 may play an important role in GDF15-induced remodeling of BM adipocytes.

**GDF15 activates the downstream genes PI3K and pAKT in bone marrow adipocyte remodeling**

Extracellular GDF15 must bind to a receptor on the membrane surface to cause intracellular changes in BM adipocytes. We screened all of the reported GDF15 receptors by RT-qPCR and found that BM adipocytes mainly express TGFβRI and TGFβRII (Figure 3A). In order to determine whether GDF15 acts through binding to TGFβRI or TGFβRII on BM adipocytes, we conducted inhibitor experiments in vitro.27,28 Western blot results showed that rhGDF15 could downregulate TRPV4 expression in BM adipocytes treated with RepSox (a TGFβRII inhibitor) rather than ITD1 (a TGFβRI inhibitor) (Figure 3B). Moreover, we found that rhGDF15 could reduce the number and area of BM adipocytes treated with RepSox as compared to those treated with ITD1 (Figure 3C, D). In accordance with data from RT-qPCR experiments, the levels of expression of lipolysis genes (ATGL and HSL) were significantly elevated in BM adipocytes treated with RepSox compared with the levels in BM adipocytes treated with ITD1 (Online Supplementary Figure S3D). These results suggest that TGFβRII is the major receptor that mediates GDF15 action on BM adipocytes.

To further verify the function of TGFβRII on BM adipocytes, we knocked down TGFβRII expression by shTGFβRII lentivirus (Online Supplementary Figure S4A, B). As shown in Figure 3E and F, when TGFβRII was knocked down in BM adipocytes, rhGDF15 did not significantly reduce the number and area of BM adipocytes. Accordingly, rhGDF15 did not significantly reduce the TRPV4 protein (Figure 3G). These results further confirmed that GDF15 regulates BM adipocyte remodeling by binding to TGFβRII.

As an important signaling downstream pathway of the TGFβ family, GDF15 could not cause significant changes in Smad2 and Smad4 proteins in BM adipocytes (Figure 3H and Online Supplementary Figure S4C). Notably, when PI3K was blocked by PI3K-IN-1 (a PI3K inhibitor), rhGDF15 could not regulate AKT phosphorylation (Figure 3I), suggesting that GDF15 is involved in the remodeling of BM adipocytes by activating the PI3K/AKT pathway.

**The PI3K/AKT pathway inhibits the TRPV4 promoter FOXC1**

PI3K/AKT acts as a signaling pathway downstream of GDF15, which may affect the transcription or translation of TRPV4. We compared the expression of different transcription factors with or without rhGDF15 treatment by RNA-sequencing analysis. The results showed that the expression of several transcription factors decreased, including FOXC1, Spalt-like gene-2 (SALL2), and MYC-associated factor X (MAX) (Figure 4A). Based on the criteria of a fold-change >2.0 and P-value <0.05, FOXC1 was identified as a transcription factor that was significantly changed in BM adipocytes after rhGDF15 treatment (Figure 4B). To investigate whether FOXC1 is responsible for GDF15 regulating TRPV4, we knocked down FOXC1 in BM adipocytes (Online Supplementary Figure S4D, E). The results showed that the expression of TRPV4 at both the mRNA and protein levels was inhibited in FOXC1 knockdown adipocytes (Figure 4C, D), suggesting that GDF15 reduced the expression of TRPV4 by negatively regulating the transcription factor FOXC1.

To further demonstrate the link between FOXC1 and TRPV4, we used a FOXC1 antibody to pull DNA fragments containing FOXC1 and used RT-qPCR to detect the TRPV4 gene in the fragment. The results showed that the control group had the sequence of the TRPV4 gene and the amount of TRPV4 gene was correspondingly decreased after knocking down FOXC1 (Figure 4E). These data suggest that FOXC1 combines directly with TRPV4. As shown in Figure 4F, rhGDF15 downregulated the expression of FOXC1 and TRPV4 protein, but PI3K-IN-1 can block this process. Taken together, these results again demonstrate that GDF15 regulates TRPV4 channels through the PI3K/AKT pathway.

**TRPV4 plays an important role in bone marrow adipocyte remodeling in acute myeloid leukemia mice**

To better understand the role of TRPV4 in leukemia cell-induced BM adipocyte remodeling, we investigated the changes in number and size of BM adipocytes with 4αPDD in mice with leukemia. FBL-3 is a mouse-derived AML cell line, which can spontaneously induce leukemia. We first confirmed that FBL-3 cells secrete GDF15 (Online Supplementary Figure S5A). In vitro, FBL-3 cells co-cultured with BM adipocytes inhibited the expression of TRPV4 protein (Online Supplementary Figure S5B). Online Supplementary Figure S5C, D shows that FBL-3 cells reduced the number and area of BM adipocytes, while 4αPDD could partly reverse this effect. Correspondingly, FBL-3 cells significantly promoted the expression of ATGL and HSL mRNA in BM adipocytes, but 4αPDD...
Figure 3. GDF15 activates the downstream genes PI3K and pAKT in bone marrow adipocyte remodeling. (A) Reverse transcriptase quantitative polymerase chain reaction (RT-qPCR) analysis of GDF15-related receptors in bone marrow (BM) adipocytes. (B) Western blot analysis of the expression of TRPV4 protein in BM adipocytes induced by recombinant human GDF15 (rhGDF15) after treatment with TGFβRI inhibitor (RepSox) or TGFβRII inhibitor (ITD1) for 4 days. (C) Oil red O staining analysis of BM adipocytes induced by rhGDF15 after treatment with RepSox and ITD1 for 6 days. All images were at a magnification of 200×. (D) The number and average area of BM adipocytes from the indicated groups were measured using Image-Pro-Plus 5.1. (E, F) BM adipocytes were infected with TGFβRII-targeted shRNA (shTGFβRII) lentivirus for 48 h and then cultured with rhGDF15 for 6 days. Adipocytes were stained with Alexa Fluor 493/503-conjugated BODIPY. 4′,6-diamidino-2-phenylindole (DAPI) stained blue and lipid droplets showed green fluorescence. The number and average area of adipocytes from the indicated groups were measured using Image-Pro-Plus 5.1. (G, H) BM adipocytes were infected with shTGFβRII lentivirus for 48 h and then cultured with rhGDF15 for 4 days. The levels of TRPV4, Smad2 and pSmad2 proteins were detected using western blot analysis. (I) BM adipocytes were treated with or without rhGDF15 and PI3K inhibitor (PI3K-IN-1, 2 μM) for 4 days. The levels of PI3K, AKT and pAKT proteins were detected using western blot analysis. β-actin protein was used as an internal control for the western blot analysis. Three independent experiments were performed. **P<0.01, *P<0.05.
could partly prevent this process (Online Supplementary Figure S5E). Additionally, 4αPDD had no significant effect on the proliferation and apoptosis of FBL-3 cells, as determined by a CCK8 assay (Online Supplementary Figure S5F) and flow cytometry analysis (Online Supplementary Figure S5G). Thus, these results suggest that this dose of 4αPDD affects adipocytes, rather than directly affecting FBL-3 cells in the co-culture system.

Figure 4. The PI3K/AKT pathway inhibits the TRPV4 promoter FOXC1. (A) Analysis of TRPV4 upstream transcription factor expression of bone marrow (BM) adipocytes treated with or without recombinant human GDF15 (rhGDF15) for 2 days by RNA sequencing. (B) Different expression of TRPV4-related transcription factor genes following rhGDF15 treatment for 2 days. (C) Reverse transcriptase quantitative polymerase chain reaction (RT-qPCR) was used to analyze TRPV4 mRNA level after treatment with FOXC1-targeted shRNA (shFOXC1) lentivirus for 48 h. (D) Western blot was used to analyze the expression of TRPV4 protein after treatment with shFOXC1 lentivirus for 48 h. (E) Chromatin immunoprecipitation-qPCR analysis of TRPV4 gene level in adipocytes with or without FOXC1 knockdown. (F) BM adipocytes were treated with or without rhGDF15 and PI3K inhibitor (PI3K-IN-1, 2 μM) for 4 days. The levels of FOXC1 and TRPV4 proteins were determined using western blot analysis. β-actin protein was used as an internal control for the western blot analysis. Three independent experiments were performed. **P<0.01, *P<0.05.
Considering that there were few adipocytes in the BM of C57BL/6 mice, we raised the mice with a high-fat diet to increase the number of adipocytes in the BM. BM adipocytes in obese mice are round or elliptical in shape (Figure 5A). Immunohistochemical staining of perilipin1 protein in the BM adipocytes showed that the BM adipocytes stained yellow (Figure 5A). According to the pathogenic characteristics of FBL-3 cells, BM samples

Figure 5. TRPV4 plays an important role in bone adipocyte remodeling in acute myeloid leukemia mice. (A) Hematoxylin & eosin staining and immunohistochemical staining with perilipin1 antibody in bone marrow (BM) sections from the controls (Ctr, n=5) and experimental mice, (mice with acute myeloid leukemia [AML], n=5; AML mice treated with 4α-phorbol 12,13-didecanoate [4αPDD], n=5). Ten fields were analyzed for each mouse at 400× magnification. Three independent experiments were performed. Scale bars represent 50 μm and 100 μm, respectively. (B, C) Adipocyte number and area in the controls, AML mice and AML mice treated with 4αPDD, quantitatively analyzed by Image-Pro-Plus 5.1. (D) Immunofluorescence was used to analyze the expression of CD117 in BM sections of the controls, AML mice and AML mice treated with 4αPDD. 4′,6-diamidino-2-phenylindole (DAPI) stained blue and CD117 showed red fluorescence. Scale bars represent 100 μm. (E) The GDF15 content of the BM supernatant in the controls and AML mice was analyzed by enzyme-linked immunosorbent assay. (F) Kaplan-Meier curves showing the overall survival rate of AML mice (n=9) and AML mice treated with 4αPDD (n=9). Three independent experiments were performed. ***P<0.001, **P<0.01, *P<0.05.
were taken 21 days after tail vein injection of FBL-3 cells. The shape of BM adipocytes did not change significantly, but the number and area decreased (Figure 5A). Further quantitative analysis showed that the number of BM adipocytes in AML mice (219±37.7/mm²) was lower than that in the controls (505±49.7/mm²) and AML mice treated with 4ePDD (334.4±39.6/mm²), but the number of BM adipocytes in AML mice treated with 4ePDD was still lower than that in the controls (the t-test for any two groups: P<0.05) (Figure 5B). Similarly, the BM adipocyte area in mice with leukemia was 860.0±142.5 µm², which was smaller than that in the controls (1686.4±106.7 µm², P<0.001). Meanwhile, the area of BM adipocytes in AML mice treated with 4ePDD was 1111.8±201.5 µm², which was larger than that of the BM adipocytes in AML mice (P<0.01), and did not return to normal (P<0.001) (Figure 5C).

We further found that there were more CD117 (a progenitor cell expression marker, red fluorescence)-positive cells in AML mice than in the AML mice treated with 4ePDD (Figure 5D). This may be due to the fact that GDF15 secreted by AML cells promotes lipolysis and is beneficial to the proliferation of leukemia cells, while 4ePDD partly prevents lipolysis. In fact, the content of GDF15 in the BM supernatant was higher in AML mice than in the controls (Figure 5E), suggesting that GDF15 secreted by leukemia cells can promote lipolysis of BM adipocytes. Furthermore, we observed that treatment with 4ePDD significantly extended overall survival of AML mice (Figure 5F). In brief, these results suggest that targeting TRPV4 in BM adipocytes can delay the progression of leukemia in mice.

Discussion

We have demonstrated a possible mechanism whereby TRPV4 mediates BM adipocyte responses to extracellular GDF15. Our data show that AML cells drive this remodeling process, at least in part, through TRPV4-dependent lipolysis in the adipocytes. Our previous reports linked increased levels of small adipocytes in BM to poor prognosis in AML patients, and revealed that GDF15 derived from leukemia cells remodels mature BM adipocytes into small adipocytes.32-33 Here, we found that GDF15 binds to its receptor TGFβRII on BM adipocytes, which in turn activates downstream target genes, including PI3K and AKT. Subsequently, TRPV4 is inhibited via downregulation of its transcription factor FOXC1. These results suggest that GDF15 regulates TRPV4 through the above pathway, thereby promoting BM adipocyte remodeling (Online Supplementary Figure S6). This finding is consistent with several reports that TRPV4 acts as a volume receptor rather than an osmotic receptor.30,32

There has been a report that TRPV4 is located on the cell membrane and acts as a calcium channel.18 Therefore, TRPV4 can regulate energy metabolism of peripheral white adipocytes by facilitating Ca²⁺ influx, which in turn stimulates the ERK1/2-dependent pathway.39 GDF15 inhibits the expression of TRPV4 in BM adipocytes, resulting in a decrease of Ca²⁺ influx (Online Supplementary Figure S3A) and an increase in pHSL protein (Online Supplementary Figure S2E) after BM adipocytes were treated with rhGDF15 for 4 days. It has been reported that reduced Ca²⁺ influx causes an increase in the expression of pHSL, leading to lipolysis of adipocytes.17 Notably, the TRPV4 channel is a tetrameric complex formed by the same or similar monomeric subunits.19 Interestingly, cytosolic N- and C-terminal domains are involved in channel gating and mediating intracellular signaling,20 indicating that it is impossible for TRPV4 to interact directly with exogenous chemical factors. Hence, it would be interesting to examine how TRPV4 communicates with extracellular GDF15.

Given our findings that extracellular GDF15 inhibited the expression of TRPV4 in BM adipocytes, we speculated that GDF15 acts on BM adipocytes through TGFβ receptors. As a member of the TGFβ superfamily, GDF15 is known to interact with receptors of TGFβ members, such as TGFβRI, TGFβRII, ALK4 and ACVR2.30-32 In addition, GDF15 has unique cognate receptors, such as GFRAL, which is mainly expressed in the central nervous system and, at low levels, in testicular tissue.30 Our data show that BM adipocytes express TGFβ receptors, but not the known unique GDF15 receptors. Moreover, TGFβRII was shown to be associated with GDF15 activity on BM adipocytes (Figure 5B-C). These experiments inform the first step of GDF15 acting on the adipocytes.

Our study further revealed that PI3K/AKT activation plays an essential role in driving GDF15 regulation of target genes in BM adipocytes. In fact, GDF15 induced the activation of Smad, a component of the classic anti-apoptosis pathway of cardiomyocytes which promotes the progression of lung cancer.60 However, we did not focus on the Smad pathway in this study because the activated Smad protein type is known to be determined by the TGFβRI present in the ligand-bound signal complex.42 In fact, GDF15 did not affect the Smad signaling pathway in BM adipocytes, which is consistent with our results (Figure 5H and Online Supplementary Figure S4C). Taken together, our data, when interpreted in the context of previous reports, suggest that the PI3K/AKT pathway may be important for GDF15-induced remodeling of BM adipocytes.

Interestingly, we observed that PI3K/AKT activation downregulated the TRPV4-associated transcription factor FOXC1. FOXC1 is also a transcription factor of ITGA7 and FGFR4 in colorectal cancer, CXCR4 in endothelial cells, and FGFR19 in ciliary body-derived cells.40-43 Moreover, the transcriptional function of FOXC1 has not been described previously for some pivotal adipogenic genes (FABP4, CEBPA and PPARC) and lipolytic genes (ATGL and HSL). Based on the knockdown of FOXC1 gene, the lipolytic gene in BM adipocytes increases (Online Supplementary Figure S4F), which confirmed that FOXC1 is important for regulating the metabolism of BM.

In addition, TRPV4 can be activated or inhibited by physical and chemical factors. When it comes to the matter of the size of cells, TRPV4 acts as a volume receptor rather than an osmotic receptor,16 suggesting that TRPV4 is involved in the regulation of cell volume. Previous studies have suggested that TRPV4 is an important inflammatory factor because TRPV4 levels are increased in inflamed tissues and activation of TRPV4 causes inflammation.44 This protein is also closely related to inflammation of white adipose tissue.45,46 However, the decreased expression of TRPV4 in leukemia-associated BM adipocytes implies that TRPV4 is not a major pro-inflammatory factor of leukemia-associated BM adipocytes. We conclude that downregulated TRPV4 preferentially pro-
motes lipolysis in BM adipocytes, contributing to their remodeling in small adipocytes in the context of AML.

Moreover, we also found that treatment with the TRPV4 agonist 4α-PDD rescued BM adipocyte remodeling, which was correlated with increased survival in AML-bearing mice, supporting a crucial role of TRPV4 in the growth and progression of AML. Although there are unstauned positive white circles in the BM of AML mice (Figure 5A), we suspect that these circles may represent an increase in blood vessels or sinuses in the BM of these animals. Leukemia cells can promote angiogenesis, which in turn contributes to the proliferation of leukemia cells. Of course, it cannot be excluded that a few adipocytes were not stained positively. Many studies have also reported the effect of circulatory factors in obese animals on leukemia. In our study, both experimental and control groups showed similar BM microenvironmental conditions, whose aging and increased adiposity of the BM microenvironment reduce the efficacy of cytotoxic chemotherapy.

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