AKAP9 Upregulation Predicts Unfavorable Prognosis in Pediatric Acute Myeloid Leukemia and Promotes Stemness Properties via the Wnt/ β -Catenin Pathway

Shiwen Wu^{1,2}
Dongqin Shen³
Li Zhao⁴

¹The First Clinical Medical College, Lanzhou University, Lanzhou, Gansu, People's Republic of China; ²Departments of Clinical Laboratory, The First Hospital of Lanzhou University, Lanzhou University, Lanzhou, Gansu, People's Republic of China; ³Department of Medical Oncology, The First Hospital of Lanzhou University, Lanzhou, Gansu, People's Republic of China; ⁴Department of Central Laboratory, Gansu Key Laboratory of Genetic Study of Hematopathy, The First Hospital of Lanzhou University, Lanzhou, Gansu, People's Republic of China

Background: PRKA kinase anchor protein 9 (AKAP9) is a scaffold protein involved in various cellular processes, including cell adhesion, proliferation, differentiation, and apoptosis. Although the oncogenic role of AKAP9 in solid tumors is well elucidated, the functions and mechanisms of AKAP9 in acute myeloid leukemia (AML) are still not understood.

Methods: We used the gene expression omnibus (GEO) database (GSE2191) to determine the mRNA expression of AKAP9 in the bone marrow of pediatric AML and healthy patients. We further used the therapeutically available research to generate effective treatments (TARGET) database to elucidate the relationship between AKAP9 expression and clinical outcomes in pediatric patients with AML. In addition, cell proliferation, cell cycle, apoptosis, RT-PCR, and Western blotting assays were applied to reveal the functions of AKAP9 and the underlying mechanisms of AKAP9 silencing in THP1 and HL60 cell lines.

Results: AKAP9 is overexpressed in the bone marrow of pediatric AML patients as compared with that of healthy patients. High expression of AKAP9 was found to be a predictor of poor overall survival (OS) and event-free survival (EFS). Using univariate and multivariate survival analyses, we found that high AKAP9 expression is an independent predictor of a worse OS and EFS. Functionally, AKAP9 silencing significantly inhibited AML cell proliferation, and cell cycle progression and promoted apoptosis. Moreover, AKAP9 silencing significantly downregulated the expression of stemness markers and β -catenin.

Conclusion: AKAP9 upregulation is a predictor of unfavorable prognosis, promotes stemness, and activates the Wnt/ β -catenin pathway in AML patients. AKAP9 may act as a prognostic biomarker of AML in pediatric patients and a future therapeutic target.

Keywords: PRKA kinase anchor protein 9, acute myeloid leukemia, prognosis, leukemic stem cells, pediatric

Correspondence: Li Zhao
Department of Central Laboratory,
Gansu Key Laboratory of Genetic Study
of Hematopathy, The First Hospital of
Lanzhou University, Lanzhou
University, No. 1 Donggang West Road,
Lanzhou, Gansu, 730000, People's
Republic of China
Tel +8613919934053
Fax +8609318356353
Email zhaoli@lzu.edu.cn

Introduction

Acute myeloid leukemia (AML) is a common hematological malignancy characterized by a clonal malignant proliferation of myeloid primordial cells.¹ AML has a poor prognosis and is the main cause of death in children and adults under 35 years of age in China.² Chemotherapy remains the primary treatment option for patients with AML. Despite the improvement in AML research and therapies,

several challenges still exist in the clinical treatment of this disease.³ Although the majority of patients achieve the stage of initial remission after induction chemotherapy, a large proportion of cases relapse and fail the re-induction chemotherapy. Accumulating evidence has suggested that leukemia stem cells (LSCs) play a key role in the recurrence and treatment failure of AML.⁴ Therefore, investigation of the mechanisms underlying the development and maintenance of LSCs and the development of more effective therapeutic targets of AML are essential.⁵

PRKA kinase anchor proteins (AKAPs) are a group of structurally diverse proteins, having the common function of binding to the regulatory subunit of protein kinase A (PKA) and limiting the holoenzyme to specific locales within cells.⁶ AKAP9 is a member of the AKAPs family, and its overexpression is implicated in solid cancer progression.⁷ But, the function of AKAP9 and its potential molecular mechanisms in hematology neoplasms, particularly AML, are unclear.

The purpose of the present study was to investigate AKAP9 expression in AML and elucidate the relationship between AKAP9 mRNA expression and the clinical prognosis of pediatric patients with AML. Additionally, we explored the function and potential molecular mechanisms involved in AKAP9-mediated progression and recurrence of AML.

Materials and Methods

Patients and Samples

The public microarray data of pediatric AML and normal control (pediatric healthy) were obtained from the gene expression omnibus (GEO) database (GSE2191) in May 2021.⁸ The differential expression of AKAP9 mRNA between 54 pediatric AML patients and 4 normal controls was analyzed by GEO2R. Moreover, we analyzed complete clinical and expression data of 285 AML pediatric patients obtained from the therapeutically available research to generate effective treatments (TARGET, <https://ocg.cancer.gov/programs/target/>) database. Patients were categorized into High-AKAP9 and Low-AKAP9 groups based on the median expression levels of AKAP9. We evaluated the correlation between mRNA expression of AKAP9 and prognosis and prognostic factors in patients with AML. The risk stratification was defined as the level of potential clinical risk based on the cytogenetic characteristics or molecular alterations as being related to those of acute myeloid leukemia.¹

Univariate and multivariate COX analyses were constructed to identify the prognostic risk factors for overall survival (OS) and event-free survival (EFS) in pediatric patients with AML.

Cell Culture

Human AML cell lines (HL-60, THP1) were obtained from the cell bank of type culture collection of the Chinese academy of sciences (Shanghai, China) and cultured under standard conditions in RPMI-1640 media (Meilunbio, Dalian, China) supplemented with 1% penicillin-streptomycin and 10% fetal calf serum. The cells were incubated at 37°C in a humidified atmosphere with 5% CO₂.

Lentivirus Infection

To determine the biological functions of AKAP9 in AML cells, we conducted AKAP9 silencing experiments. AML cells were infected with lentivirus containing either a short hairpin RNA for AKAP9 (sh-AKAP9) or a negative control vector (sh-NC) according to the manufacturer's instructions. We observed green fluorescence at 48 h after lentivirus transfection using a fluorescence microscope (magnification, 100x). Puromycin was used for screening of the infected cells at 72 h after transfection. The silencing efficiencies of AKAP9 were identified by real-time quantitative polymerase chain reaction (RT-PCR) and Western blot analysis. shRNAs were synthesized by Genechem (Shanghai, China). The sequence of the AKAP9 shRNA used was 5'-GCCAAGAAGAAAGATTGATTAAATCAATCTTTCTTCTTGCC-3'.

RNA Isolation and Real-Time

Quantitative Polymerase Chain Reaction

The total RNA was extracted using TRIzol reagent. The target cDNA was obtained by reverse transcription using a FastKing gDNA Dispelling RT SuperMix kit (TIANGEN, Beijing, China) according to the manufacturer's instructions. A SuperReal PreMix Plus kit (TIANGEN, Beijing, China) was used to perform RT-PCR with specific primers as previously described.^{7,9} Primers for RT-PCR were synthesized by TaKaRa Biotechnology (Dalian, China). Each sample was tested on triplicates, and the mRNA expression was measured by the threshold cycle (CT) values. The relative expression of

mRNA was calculated by the $2^{-\Delta\Delta CT}$ method, and β -actin was used as an internal reference for mRNAs.

Protein Isolation and Western Blot Analysis

Total protein was extracted from the cells with RIPA buffer. The concentration of protein was determined by the BCA protein assay kit (MeiLunBio, Dalian, China). Equal amounts of protein were separated by SDS-PAGE and transferred onto the polyvinyl difluoride (PVDF) membranes. The membranes were blocked for 2 h with 5% non-fat milk in tris-buffered saline (TBS) at room temperature and incubated with primary antibodies against AKAP9 (1:500 dilution, Abcam, catalog ab32679), β -catenin (1:1000 dilution, Proteintech, catalog 66379-1-Ig), CD133 (1:1000 dilution, Proteintech, 18470-1-AP), ALDH1A1 (1:1000 dilution, Proteintech, 15910-1-AP), and β -actin (1:1000 dilution, Proteintech, catalog 66009-1-Ig) at 4°C overnight. The membranes were then washed with TBS-tween-20 thrice and incubated with anti-mouse or anti-rabbit secondary antibodies (1:5000 dilution, Proteintech, catalog SA00001-1, and SA00001-2) for 2 h at room temperature. The protein bands were treated with ECL reagent (Beyotime Biotechnology, Shanghai, China) and visualized by Amersham Imager 600 visualizer (GE Healthcare, Pittsburgh, USA). The protein expression level was quantified and analyzed using ImageJ software (National Institutes of Health) software.

Cell Proliferation Assays

Cell proliferation ability was evaluated using cell counting kit-8 (CCK8) (MeiLunBio, Dalian, China) according to the manufacturer's protocol. First, the cells were inoculated into 96-well plates at a density of 5×10^4 /well and treated with CCK-8 solution at 0, 24, 48, 72, and 96 h. Finally, the absorbance of each well was measured by a microplate spectrophotometer at 450 nm.

Cell Cycle Assays

For cell-cycle analysis, the cells were washed with cold PBS, and then fixed with 70% ethanol at 4°C overnight. The cells were washed twice with PBS and incubated with 1 mg/mL propidium iodide and 0.5 mg/mL RNase A (MultiSciences Biotech, Co., Ltd., Hangzhou, China) for 30 min in dark at 37°C. The phase change of the cell cycle was examined using a flow cytometer (BD Biosciences, San Jose, CA, USA). The flow cytometry

data were analyzed using ModFit LT software for Windows (Version 5.0.9). All experiments were conducted in independent triplicates.

Cell Apoptosis Analysis

Annexin V-phycoerythrin (APC) and 7-amino-actinomycin D (7-AAD) apoptosis detection kits (MultiSciences Biotech, Co., Ltd., Hangzhou, China) were used to detect the apoptosis of THP1 and HL60 cells after AKAP9 silencing according to the manufacturer's protocols. In short, 5×10^5 cells were washed twice with cold PBS and resuspended in 500 μ L binding buffer. Next, the cells were stained with 5 μ L Annexin V-APC and 5 μ L 7-AAD for 15 min in the dark at room temperature. The apoptotic cell percentage was calculated using a flow cytometer (BD Biosciences, San Jose, CA, USA). The flow cytometry data were analyzed by FlowJo (version 10.7.1) software.

Statistical Analysis

Statistical analyses were carried out using SPSS 22.0 software (IBM, Armonk, New York, USA). OS and EFS were evaluated by the Kaplan–Meier method and analyzed using the Log rank test. Univariate and multivariate analyses of prognostic factors in patients with AML were conducted using the Cox regression model. Multivariate analysis included variables with $P < 0.1$ in univariate analysis. Pearson Chi-square and Fisher's exact tests were used for categorical variables. Quantitative variables were expressed as mean $\pm$ SD. The Student's *t*-test was used to determine differences between the two groups. A *P*-value less than 0.05 was considered statistically significant.

Results

AKAP9 Expression in Pediatric AML

To explore the role of AKAP9 in AML pathogenesis, we evaluated the expression of AKAP9 in AML patients and normal control based on the GSE2191 from the GEO database, which contained 4 normal controls and 54 pediatric patients with AML. Differences in AKAP9 expression levels between pediatric AML and the normal group were evaluated using the Mann–Whitney *U*-test. The findings revealed that AKAP9 has a significantly higher expression in pediatric patients with AML compared to normal control ($P < 0.01$) (Figure 1A).

To explore whether AKAP9 expression might play a role in relapse and drug resistance of AML in pediatric patients,

Figure 1 The AKAP9 mRNA expression and prognosis in pediatric patients with AML. **(A)** The AKAP9 mRNA expression level in pediatric AML and control in the GSE2191 cohort. **(B)** Patients with a high AKAP9 expression had a significantly higher MRD positive rate when compared with those with a low AKAP9 expression. **(C)** Patients with a high AKAP9 expression showed significantly higher CR rate when compared with those with a low AKAP9 expression. **(D and E)** High AKAP9 mRNA expression proved to be a predictor of poor overall survival and event-free survival, as ascertained from the TARGET database. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

we compared the positive rate of minimal residual disease (MRD) and complete remission (CR) rate between high- and low-AKAP9-expression, respectively, at end of the first and second course of chemotherapy. We found that the MRD rate in the high-AKAP9 group was significantly higher than that in the low-AKAP9 group at end of the first course (37.1 vs 14.8%; $P < 0.01$) and second course (20.3 vs 7.7%; $P < 0.01$) of chemotherapy (Figure 1B). The CR rate in the low-AKAP9 group was significantly higher than that in the high-AKAP9 group (88.7% vs 65.7%, $P < 0.01$) at end of the first course of chemotherapy; however, no significant differences were observed at the end of the second course of chemotherapy in patients with low and high AKAP9 expression (88.0 vs 81.8%; $P=0.163$) (Figure 1C).

AKAP9 Overexpression is a Predictor of Poor Prognosis in Pediatric AML Patients

To determine the applicability of AKAP9 in the prognosis of pediatric AML, we analyzed the relationship between

the AKAP9 expression level and OS/EFS time based on the TARGET database. We found that patients with high AKAP9 expression had a shorter OS and EFS time compared to patients with low AKAP9 expression (Figure 1D and E). These findings suggest that the high expression of AKAP9 can result in poor survival in pediatric patients with AML. Therefore, AKAP9 can act as a significant biomarker to determine the prognosis of pediatric AML patients.

Association Between AKAP9 Expression and Clinical Characteristics of Pediatric AML

We investigated the association between the AKAP9 expression and the clinicopathological features of pediatric patients with AML. Patients with low AKAP9 expression showed higher white blood cell counts than those with high AKAP9 expression (median, $59.0 \times 10^9/L$ vs $31.1 \times 10^9/L$; $P < 0.01$). Significant differences were observed in

Table 1 Comparison of Clinical Manifestations and Laboratory Features Between Low-AKAP9 and High-AKAP9 AML Patients

Characteristics	LowAKAP9 (n=142)	HighAKAP9 (n=143)	P
Age/years, median (range)	11(0–22)	9(0–23)	0.202
Gender, no. (%)			0.438
Male	79(55.63)	73(51.05)	
Female	63(44.37)	70(48.95)	
Race, no. (%)			0.840
Caucasian	105(73.94)	105(73.43)	
African American	16(11.27)	16(11.19)	
Asian	3(2.11)	6(4.20)	
Other	6(4.23)	7(4.90)	
Unknown	12(8.45)	9(6.30)	
WBC at diagnosis/ ×10 ⁹ /L, median (range)	59.0(1.6–446)	31.1(0.9–519)	<0.01
BM blast/ %, median (range)	77.5(31–100)	72.0(14–100)	0.009
PB blast/ %, median (range)	66.0(0–97)	52.0(0–97)	<0.01
Risk stratification, no. (%)			0.044
Low	67(47.18)	47(32.87)	
Standard	51(35.92)	74(51.75)	
High	18(12.68)	15(10.49)	
Unknown	6(4.23)	7(4.90)	
Karyotype, no. (%)			0.677
Normal	34(23.94)	37(25.87)	
Abnormal	101(71.13)	96(67.13)	
Unknown	7(4.93)	10(6.99)	
FLT3-ITD, no. (%)			0.894
Positive	23(16.20)	24(16.78)	
Negative	119(83.80)	119(83.22)	

Abbreviations: WBC, white blood cells; BM, bone marrow; PB, peripheral blood; FLT3-ITD, internal tandem duplication of the FLT3 gene.

the bone marrow (BM) blast percentages ($P = 0.009$), peripheral blood (PB) blast percentages ($P < 0.01$), and risk stratification ($P = 0.044$) between the two groups. No significant relationships were noted between the AKAP9 expression and age, gender, race, karyotype status, and internal tandem duplication of the *FLT3* (FLT3-ITD) (Table 1).

AKAP9 Overexpression is Independently Associated with Poor Prognosis of Pediatric AML Patients

To investigate whether AKAP9 is an independent prognostic factor in AML, we conducted univariate and multivariate Cox analysis for OS and EFS using the TARGET

database. The univariate analysis showed that three factors – FLT3-ITD, risk stratification, and AKAP9 expression – are risk markers associated with shorter OS, while four factors – FLT3-ITD, risk stratification, FAB category, and AKAP9 expression – are risk markers associated with shorter EFS. As per the findings of multivariate analysis, AKAP9 overexpression is an independent prognostic factor for poor OS and EFS in pediatric AML patients (Figure 2A and B).

Establishment of AKAP9 Stable Down-Regulation AML Cell Line

To investigate the biological functions of AKAP9 in AML, THP1, and HL60 cells were infected with lentivirus containing AKAP9-shRNA to conduct stable knock-down of AKAP9. As shown in Figure 3A, the cells that were successfully transfected with lentivirus showed green fluorescence after 48 h. The RT-PCR and Western blot results revealed that the mRNA and protein levels of AKAP9 in THP1 and HL60 cells infected with lentivirus containing AKAP9-shRNA were significantly downregulated as compared with the corresponding levels in the control cells ($P < 0.001$; Figure 3B–D).

AKAP9 Silencing Inhibits Cell Proliferation and Cell Cycle Progression of AML Cells

CCK8 assays were performed to evaluate the effect of AKAP9 silencing on cell proliferation in AML cells. According to the growth curve, the proliferation of THP1 cells and HL60 cells was significantly suppressed after AKAP9 silencing, while cells transfected with shNC exhibited no significant inhibition of cell proliferation (Figure 4A and B).

To explore whether AKAP9 is involved in the cell cycle of AML cells, we assessed the effect of AKAP9 silencing on the cell cycle distribution of THP1 cells and HL60 cells using FACS. The findings suggested that AKAP9 silencing slightly, but statistically significantly, prolonged the G0/G1 phase (5.38% and 3.87%) and reduced the G2/M phase (4.14% and 4.91%) in both THP1 and HL60 cells; however, the S phase was not significantly changed. As expected, the G0/G1 phase, S phase, and G2/M phase were not changed in sh-NC transfected cells (Figure 4C–F). Together, these findings revealed that AKAP9 gene silencing significantly decreases the cell proliferation potential and significantly inhibits the cell cycle progression of AML cells.

Figure 2 Univariate and multivariate COX analyses of prognostic factors for OS and EFS in pediatric patients with AML. **(A)** The AKAP9 overexpression is an independent prognostic factor of worse OS in pediatric patients with AML. **(B)** AKAP9 overexpression is an independent prognostic factor of worse EFS in pediatric patients with AML. **Abbreviations:** CI, confidence interval; HR, hazard ratio.

Figure 3 Establishment of AKAP9-silenced THP1 and HL60 cell lines. **(A)** Fluorescence images of THP1 and HL60 cells at 48 h after transfection. **(B)** The mRNA and **(C)** protein expression levels of AKAP9 in THP1 and HL60 were reduced in AKAP9 silenced-cell lines. **(D)** Quantitative analysis of the AKAP9 protein expression in **(C)**. **** $P < 0.0001$.

AKAP9 Silencing Promotes the Apoptosis of AML Cells

We determined whether AKAP9 is involved in AML cell apoptosis. Annexin V APC/7-AAD double-staining assays revealed that the average percentage of apoptotic cells in the sh-AKAP9 group (THP1, 11.23%; HL60, 11.61%) was significantly higher than the corresponding percentage in sh-NC (THP1, 7.15%; HL60, 5.52%) and control (THP1, 7.52%; HL60, 5.80%) groups ($P < 0.01$, **Figure 5A** and **B**). There were no significant differences in the apoptotic rates between the control and sh-NC groups ($P > 0.05$; **Figure 5B**). The results revealed that

AKAP9 silencing significantly promotes apoptosis in THP1 and HL60 cells.

AKAP9 Silencing Decreased Stem Cell Marker Expression of AML Cells

To further investigate the effect of AKAP9 on the stemness of AML cells, we analyzed the stem cell biomarkers including ALDH1A1 and CD133 by Western blot. The expression levels of ALDH1A1 in THP1 and HL60 cells were significantly decreased after AKAP9 silencing (**Figure 6A–C**). These data suggest that AKAP9 might be a regulator of AML stem cell phenotype.

Figure 4 Effects of AKAP9 silencing on the proliferation and cell cycle of THP1 and HL60 cells. (A and B) AKAP9 silencing decreased the potential of THP1 and HL60 cells' proliferation. (C and D) and (E and F) AKAP9 silencing inhibited the cell cycle progression of THP1 and HL60 cells. * $P < 0.05$, ** $P < 0.01$.

Figure 5 AKAP9 silencing promoted the apoptosis of HL60 and THP1 cells. (A) Apoptotic cells were analyzed by flow cytometry by staining annexin APC/7-AAD. (B) The percentage of apoptosis in THP1 and HL60 cells after AKAP9 silencing. ** $P < 0.01$, *** $P < 0.001$.

AKAP9 Silencing Suppresses the Wnt/ β -catenin Signaling Pathway in AML Cells

The Wnt/ β -catenin signaling pathway is commonly involved in the stemness of cancer cells. The inhibition of Wnt/ β -catenin signaling can reduce the stemness of AML cells. To determine whether AKAP9 participates in regulating the stemness of AML cells via Wnt/ β -Catenin signaling, we evaluated the relationship between AKAP9 and β -catenin (CTNNB1), which is a key regulatory molecule of the Wnt signaling pathway. The mRNA expression level of CTNNB1 in AKAP9-high patients was significantly higher than that in AKAP9-low patients (Figure 7A). We also observed a significant correlation between AKAP9 and β -catenin mRNA expressions ($P < 0.001$) (Figure 7B). Furthermore, Western blot assays were conducted to determine the expression of AKAP9 and intracellular signal transducer β -catenin in AKAP9-modulating cells. The results showed that AKAP9 down-regulation suppressed the expression of β -catenin (Figure 7C and D). All these findings suggested that AKAP9 silencing inhibits the Wnt/ β -Catenin signaling pathway in AML cells.

Discussion

AKAP9, also known as Yotiao, is linked to the development and metastasis of several human cancers,

including lung cancer,¹⁰ breast cancer,^{11,12} colorectal cancer,^{7,13} melanomas,¹⁴ thyroid carcinomas,^{15,16} bladder cancer,¹⁷ and oral cancer.¹⁸ However, the functions and mechanisms of AKAP9 in these tumors are not yet completely elucidated. Also, there are no available demonstrating the role of AKAP9 in the development of AML. Our results suggested that AKAP9 expression is upregulated in pediatric AML patients, and its high expression is associated with poor prognosis. AKAP9 overexpression was found to be independently associated with poor prognosis of pediatric AML patients. AKAP9 plays an important role in the progression and relapse of AML; however, the related underlying mechanisms remain poorly understood.

LSCs are the subset of cells exhibiting quiescent and low energy production state and possess limitless self-renewal properties.¹⁹ The development of treatment strategies targeting LSCs is expected to facilitate the development of a more effective cure for AML. It is unclear whether AKAP9 plays a critical role in LSCs. AKAP9 is the key regulator in the G1 to S phase transition.²⁰ Our cell experiments confirmed that AKAP9 silencing could inhibit the growth of AML cells and regulate the cell cycle progression. MRD is a significant predictor of AML recurrence and prognosis. MRD positive status

Figure 6 Effect of AKAP9 silencing on the stem cell-related molecular markers in HL60 and THP1 cells. **(A)** Western blotting of stem cell-related molecular markers—ALDH1A1 and CD133—in the control (Ctrl), sh-NC-, and sh-AKAP9-transfected THP1 and HL60 cells. **(B and C)** Densitometry analyses of the AKAP9 protein expression levels in THP1 and HL60. *** $P < 0.001$.

Figure 7 AKAP9 silencing suppressed the Wnt/ β -catenin signaling pathway in AML cells. **(A)** The mRNA expression level of CTNNB1 in AKAP9-high and AKAP9-low patients from the TARGET database. **(B)** The correlation of CTNNB1 and AKAP9 mRNA expressions in pediatric AML patients from the TARGET database. **(C)** Western blotting of β -catenin in the control (Ctrl), sh-NC-, and sh-AKAP9-transfected THP1 and HL60 cells. **(D)** Densitometry analysis of the β -catenin protein expression levels in **(C)**. *** $P < 0.001$, **** $P < 0.0001$.

indicated that a small amount of LSC remained in the bone marrow protective environment even after the complete remission of chemotherapy (or stem cell transplantation).^{21,22} Patients with high AKAP9 expression had a higher positivity rate of MRD compared with patients with low AKAP9 expression. Furthermore, ALDH1A1 and CD133 are important markers in LSCs.^{23,24} ALDH1A1 plays an important role in initiation, relapse, chemotherapy resistance, and progression of AML. These processes are involved in the prognosis of AML.^{25–27} Moreover, CD133 expression is highly associated with chemoresistance and prognosis of patients with AML.²⁸ Our results showed that AKAP9 silencing decreases the expression of ALDH1A1 and CD133 in AML cells. The findings support the hypothesis that AKAP9 overexpression contributes to AML cell stemness.

The Wnt/ β -catenin pathway is at the center of carcinogenesis and cancer progression. It is necessary for the maintenance and self-renewal of LSCs. β -catenin is considered a vital downstream effector of the Wnt/ β -catenin pathway.^{29,30} Several molecules involved in the Wnt/ β -

catenin pathway have a prognostic value in AML.²⁹ Therapeutics targeting the Wnt/ β -catenin pathway have a great potential to treat AML. Studies have shown that the AKAP gene family is involved in the Wnt/ β -catenin pathway. The AKAP family members can not only bind to β -catenin but also directly interact with GSK3 β , which is required for the regulation of β -catenin and thus for Wnt signaling.^{31–33} Our results reveal that AKAP9 and β -catenin mRNA expressions are highly correlated, and AKAP9 silencing suppresses the protein levels of β -catenin, indicating that AKAP9 may promote the activation of β -catenin and Wnt/ β -catenin signaling pathway.

Conclusion

In conclusion, AKAP9 mRNA expression is upregulated in pediatric AML patients, and its expression influences patient prognosis. AKAP9 silencing significantly inhibits AML cell proliferation and cell cycle progression and promotes apoptosis. Moreover, AKAP9 increases LSCs stemness by activating the Wnt/ β -catenin signaling pathway. AKAP9 may serve as a prognostic indicator for pediatric AML and provide a novel molecular drug target

for AML. Further in vivo experiments and patient-derived sample studies are needed to validate our findings.

Abbreviations

AKAP9, A-kinase anchor proteins 9; FAB category, French-American-British classification category; FLT3-ITD, internal tandem duplication of the FLT3 gene; EFS, event-free survival; OS, overall survival; TARGET, the therapeutically available research to generate effective treatments database; GEO, the gene expression omnibus database.

Ethical Statement

The present study was approved by the Ethics Committee of LZU No. 1 Hospital.

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Disclosure

The authors report no conflicts of interest in this work.

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