Clinical profile in KMT2A-SEPT6-positive acute myeloid leukemia: Does it often co-occur with NRAS mutations?

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Background: The KMT2A-SEPT6 fusion gene is a relatively rare genetic event in leukemia. Its clinical characteristics and prognosis, especially the profile of co-occurring gene mutations remain unclear.

Methods: We retrospectively analyzed the characteristics of four cases carrying KMT2A-SEPT6 in our hospital, and provided a literature review.

Results: All the four patients were diagnosed with acute myeloid leukemia (AML) and harbored X chromosome and 1 chromosome rearrangements, they all manifested high levels of D-dimer. Three of four patients had NRAS mutations while one patient with congenital AML did not. Of the four cases, one developed drug resistance, one suffered relapse after bone marrow transplantation (BMT) and two died. Combined with other cases reported in the literature, we found that of all patients diagnosed with AML, 90.9% were children (≤9 years old). Patients with white blood cells ≥20.0 × 10⁹/L or diagnosed with M4 had a shorter overall survival (P < 0.05). Age, whether to receive BMT, and the chromosome rearrangement patterns had no significant effect on overall survival (P > 0.05).

Conclusions: KMT2A-SEPT6 was more commonly observed in pediatric AML patients, some of which may co-occur with NRAS mutations. The prognosis was related to the white blood cell levels and the leukemia subtype, but was not related to age or BMT. More cases need to be accumulated to better understand the profile in KMT2A-SEPT6-positive AML.

KEYWORDS
acute myeloid leukemia, gene rearrangement, KMT2A-SEPT6, mutations, NRAS

Introduction

Acute myeloid leukemia (AML) is a malignant tumor that originates from myeloid blood cells and is characterized by abnormally increased leukemic cells in bone marrow or peripheral blood (1). Cytogenetic and molecular abnormalities commonly occur in AML patients. Based on genetic mutations and specific chromosomal rearrangements, the World Health Organization (WHO) divides AML with recurrent genetic abnormalities into different subgroups (2). Translocations involving the lysine
(K)-specific methyltransferase 2A gene (KMT2A), previously known as mixed lineage leukemia (MLL), are common chromosomal abnormalities in AML (3, 4). The KMT2A gene at 11q23 has many partner genes, of which over 80 have been identified (5).

Among them, SEPT6 located at Xq24 which is involved in the formation of the KMT2A arrangement t(X;11)(q22-24;q23), is extremely rare in AML (6). SEPT6 is a member of Septins family, an evolutionarily conserved family of GTP-binding proteins that associate with cell polarity, cytokinesis and oncogenesis (7, 8). Five septin genes (SEPT2, SEPT5, SEPT6, SEPT9, and SEPT11) were identified as KMT2A fusion partner (9). SEPT6, which plays a role in actin and microtubule cytoskeletons, is involved in infectious diseases, Down’s syndrome, schizophrenia, bipolar disorder and cancers, including leukemia and lymphoma (10, 11). To our knowledge, only a limited number of KMT2A-SEPT6-positive cases have been documented in the literature. Most cases are children while only one adult case has been reported (6, 12–21). The exact role of KMT2A-SEPT6 in hematopoietic cells and its effect on leukemogenesis are still unknown. There is little information on the clinical features, treatment strategies and prognosis of such patients, and accompanying gene mutations carried by such patients have not been described.

We retrospectively analyzed data from four acute leukemia patients carrying the KMT2A-SEPT6 fusion gene who were treated in our hospital, especially gene mutation information. Additionally, we reviewed cases in the literature together with our cases to provide evidence for potential therapeutic strategies.

**Materials and methods**

**Case selection**

We collected four patients harboring the KMT2A-SEPT6 gene from a pool of 1,656 leukemia patients within our hematological disease database in the past 4 years, which were referred to as cases 1–4. The diagnostic criteria were according to the WHO classification of tumors of hematopoietic and lymphoid tissues (22). We conducted a retrospective analysis and systematic summary with information on morphology, flow cytometric analysis, cytogenetics, molecular biology, and other related laboratory test results.

**Literature review**

We conducted a literature search on PubMed with the keywords “KMT2A-SEPT6,” “MLL-SEPT6,” “KMT2A-SEPTIN6,” “MLL-SEPTIN6” or “t(X;11)” to gather related case reports.

**Statistical analysis**

Kaplan–Meier method was used for survival evaluation. The log-rank test was used to assess the difference between groups. \( P < 0.05 \) was considered statistically significant. All data were analyzed with SPSS Statistics, version 21 (StatSoft).

**Results**

**Clinical presentation**

All KMT2A-SEPT6-positive cases (cases 1–4) were male with ages ranging from 0 to 57 years, and their detailed information is summarized in Table 1. All cases had manifestations of fever and pale complexion. The three pediatric cases (cases 2, 3, and 4) were accompanied by hepatosplenomegaly, and two cases (cases 3 and 4) had scattered petechiae and ecchymoses. Case 1 was an elderly male patient with perianal abscess and diabetes in addition to the above symptoms. Case 2 was accompanied by pain in both lower limbs, tenderness of the sternum and hypertrophy of the tonsils. In case 3, multiple lymph nodes were palpable on the bilateral neck, he was positive for sternal tenderness, and he also had hyperuricemia and acute bronchial pneumonia. Case 4 was a newborn delivered by cesarean section due to a “decreased fetal heart rate.” The birth weight was 3,100 g. The infant had no spontaneous breath at birth and was generally cyanotic. He restored spontaneous respiration under assisted ventilation, and he developed persistent pulmonary arterial hypertension and neonatal pneumonia. His parents were healthy and had no history of genetic disease.

Laboratory results showed that three cases (cases 1, 3, and 4) had high white blood cell (WBC) counts, anemia, and low platelet counts. One patient (case 2) had low hemoglobin levels and WBC counts but normal platelet counts, and two patients (cases 2 and 3) had increased serum lactate dehydrogenase levels. The D-dimer levels of all four patients were increased.

**Morphological evaluation**

All cases exhibited morphological characteristics of AML (Figure 1). Leukemia cells have the morphological characteristics of blast cells, including medium to large nuclei with round and often eccentric nuclei, loose chromatin, abundant basophilic cytoplasm, and Auer bodies can be seen in some cells. The French American British (FAB) morphological classification of each case was M5 (cases 1 and 4), M2 (case 2) or M4 (case 3). Three cases (cases 1, 3, and 4) showed hypercellular bone marrow, and the other case (case 2) had severe hypocellular bone marrow. Case 1 and case 4 were evaluated as M5 and revealed 92.0 and 20.4% blasts in marrow aspirate and 69.0 and 17.0% blasts in peripheral blood, respectively (Table 1;
TABLE 1  Clinical features of four KMT2A-SEPT6-positive AML patients.

| Items                          | Case 1                  | Case 2                  | Case 3                  | Case 4                  |
|--------------------------------|-------------------------|-------------------------|-------------------------|-------------------------|
| **Sex**                        | Male                    | Male                    | Male                    | Male                    |
| **Age**                        | 57 years                | 9 years                 | 16 months               | 0 month                 |
| **Physical examination**       | Perianal abscess        | Lower limbs pain, sternal tenderness, and tonsil hypertrophy | Scattered petechiae on the neck, cervical lymphadenopathy and sternal tenderness (+) | Scattered petechiae and ecchymoses |
| **Hematomegaly/splenomegaly**  | No/No                   | Yes/Yes                 | Yes/Yes                 | Yes/Yes                 |
| **WBC/Hb/PLT (×10⁹/L/g/L/×10⁹/L)** | 12.3/79.0/105.0         | 3.0/93.0/245.0          | 123.8/41.0/39.0         | 112.0/90.0/9.0          |
| **Serum LDH (U/L)**            | NA                      | 966                     | 2,080                   | NA                      |
| **D-Dimer [µg/L(DDU)]**        | 13117.0                 | 3149.0                  | 2469.0                  | 620.0                   |
| **Blood blasts (%)**           | 69.0                    | 2.0                     | 52.0                    | 17.0                    |
| **Bone marrow blasts (%)**     | 92.0                    | 27.5                    | 56.0                    | 28.4                    |
| **Morphological diagnosis**    | AML-M5                  | AML-M2                  | AML-M4                  | AML-M5                  |
| **Immunophenotype**            | The leukemic cells      | The leukemic cells      | The leukemic cells      | The leukemic cells      |
|                                | expressed HLA-DR,       | expressed HLA-DR,       | expressed HLA-DR,       | expressed HLA-DR,       |
|                                | CD117, CD33, CD13,     | CD33, CD38, CD15,      | CD33, CD38, CD15,      | CD33, CD38, CD15,      |
|                                | CD38, CD15, CD64, and  | CD64, and CD4           | CD15, and CD64          | CD64                    |
|                                | CD4                     |                         |                         |                         |
| **Karyotype**                  | 46, Y, t(X;11)[q24q23][4]/46,XY[6] | 45, Y,del(X)(q21),der(11) | 46, Y,t(X;11)[q24q23],[20,add(22)[q13][3] | 46,Y,ins(X;11)          |
|                                |                         |                         |                         |                         |
| **Gene mutations**             | NRAS:                   | NRAS:                   | NRAS:NM_002524:exon4.c. | Negative                |
|                                | NM_002524:exon2.c.G3ST: | NM_002524:exon2.c.     | G36A:p.A146T, VAF = 0.44, Missense |
|                                | p.G12V, VAF = 0.04, Missense; ASXL1: | G3ST:p.G12V, VAF = 0.03, Missense |                         |                         |
|                                | NM_015338:exon12.c.2464 | dup:A:p.D8216, VAF = 0.36, Frameshift |                         |                         |
| **Treatment protocol**         | IA regimen (resistance); then changed to decitabine combined with half-dose CAG regimen (once) + high dose cytarabine | MAE regimen + intrathecal chemotherapy + BMT | HA regimen               | No chemotherapy         |
| **CR**                         | Yes                     | Yes                     | Yes                     | NA                      |
| **Relapse**                    | Yes                     | Yes                     | Yes                     | NA                      |
| **Follow-up (months)**         | 18                      | 43                      | 36                      | 0.5                     |
| **Outcome**                    | Alive                   | Alive                   | Died                    | Died                    |

CNS, central nervous system; WBC, white blood cells; Hb, hemoglobin; PLT, platelets; LDH, lactate dehydrogenase; VAF, variant allele frequencies; CAG, cytarabine + adacinomycin + granulocyte colony stimulating factor; MAE, Mitoxantrone + cytarabine + etoposide; BMT, bone marrow transplantation; HA, homoharringtonine + cytarabine; CR, complete remission; NA, not available.

**Figures 1A,D)**. Case 2 was evaluated as M2; the marrow aspirate revealed 27.5% myeloblasts and the peripheral blood exhibited 2.0% blasts (Table 1; Figure 1B). Case 3 was evaluated as M4; the marrow aspirate showed 56.0% blasts and the peripheral blood exhibited 52.0% blasts (Table 1; Figure 1C).

**Flow cytometric analysis**

Flow cytometric analysis revealed the presence of myeloid blasts in bone marrow samples from all four patients (Table 1; Figure 2). The percentage of...
Morphologic evaluation of leukemic cells at diagnosis (Wright–Giemsa stain, ×1,000). (A–D) represent cases 1, 2, 3, and 4, respectively. BM, bone marrow; PB, peripheral blood.

The sequencing depth was 2,000×. The results showed that three cases (cases 1–3) harbored NRAS (NM_002524) mutations, and the mutation sites of cases 1 and 2 were both NRAS G12V, with variant allele frequencies (VAFs) of 0.04 and 0.03, respectively. In addition, case 1 was also accompanied by an insertion mutation of ASXL1 D821 with a VAF of 0.36. The reexamination results at the third month showed that the KMT2A-SEPT6 gene remained positive but its expression level dropped from 100 to 24.4%. The ASXL1 D821 VAF dropped to 0.01, and the NRAS G12V gene mutation was not detected. Four months later, the patient achieved complete remission (CR), the ASXL1 D821 VAF dropped to 0.003, and both the NRAS mutation and KMT2A-SEPT6 fusion gene were negative. The NRAS A146T mutation in case 3 occurred in exon 4 with a VAF of 0.44. KMT2A-SEPT6 gene and NRAS gene mutations both turned negative on the ninth month after diagnosis. The patient relapsed on the twenty-seventh month with positive KMT2A-SEPT6 and NRAS A146T mutation (VAF = 0.27). In case 4, no pathogenic gene mutations were detected.

Clinical course

The treatment and follow-up information of all cases (cases 1–4) is shown in Table 1. Three patients (cases 1–3) received chemotherapy, patient 2 subsequently received a bone marrow transplantation (BMT), and patient 4 was a newborn and did not receive any chemotherapy. The clinical follow-up period ranged from 0.5 to 43 months, with a median of 27 months. Case 1 received induction chemotherapy of IA (idarubicin + standard cytarabine) regimen, and revealed non-remission with blasts from 92.0 to 40.5% in bone marrow. Then, he received D-CAG (decitabine combined with low-dose cytarabine + aclacinomycin + granulocyte colony stimulating factor) and achieved CR. The KMT2A-SEPT6 gene expression reduced from 100 to 24.4%, and the NRAS G12V gene mutation was negative after the second chemotherapy. Subsequently, he received high-dose cytarabine consolidation therapy and an intrathecal injection (cytarabine + dexamethasone + methotrexate) for central nervous system infiltration prevention. The patient revealed a normal karyotype and negative molecular results of KMT2A-SEPT6 rearrangement and NRAS mutation, and sustained CR for 10 months. However, he relapsed 3 months later. From this time point the patients

Blasts was highest at 92% in case 1 and lowest at 4.2% in case 2. All cases were positive for CD33, CD15, and CD64, indicating myeloid lineage, and CD13 was positive only in case 1 and case 3. All the cases were positive for CD38 and HLA-DR. CD117 was positive only in case 1 and case 3. The results showed that all four cases had clonal abnormalities of the X chromosome and chromosome 11 or complex karyotype abnormalities (Table 1; Figure 3). In case 1, the metaphase cells exhibited abnormalities of t(X;11)(q24;q23) (Figure 3A). The metaphase cells collected in case 2 showed 45, Y, del(X)(q21), der(11)(X;11)(q24;q23), −20, add (22)(q13) (Figure 3B). In case 3, in addition to the abnormal karyotype of t(X;11)(q24;q23), thirteen metaphase cells had del(7)(q21q31) (Figure 3C). The metaphase cells in case 4 showed an abnormal karyotype of 46, Y, ins(X;11)(q23;q24q12) (Figure 3D).
had a stable blast count of 5.0%-10.5% to the last follow-up.

Case 2 received MAE (mitoxantrone + standard cytarabine + etoposide) regimen and reached CR. Then he received high-dose cytarabine consolidation therapy. After that, the patient received allogeneic hematopoietic stem cell transplantation (allo-HSCT). The conditioning regime was modified BU/CY + ATG (busulfan/cyclophosphamide + antithymocyte globulin, peking regime). The donor was his father, HLA 5/10, A+ to B+. We used CsA + MTX + MMF (cyclosporine + methotrexate + mycophenolate mofetil) for Graft-Versus-Host Disease (GVHD) prophylactic. The process of transplantation was well-off, and bone marrow assessment results were negative continuously. However, 19 months after transplantation, he relapsed with central nervous system leukemia. Following five courses of intrathecal injection, the child achieved CR again.

Case 3 received HA (homoharringtonine + standard cytarabine) regimen and intrathecal injection (cytarabine + dexamethasone + methotrexate). He reached CR 4 weeks after diagnosis.
patient relapsed 27 months later and died 36 months after diagnosis. Case 4 had dyspnea at birth, and he was on assisted ventilation and given blood infusion to improve anemia and thrombocytopenia. The newborn’s condition did not improve during the treatment, and the child died 2 weeks later.

**Literature review**

A total of 22 KMT2A-SEPT6-positive cases were included in this literature review, including four cases in our report and eighteen cases from the literature (6, 12–21). Tables 2, 3 list the detailed clinical information of all patients and clinical features of evaluable patients.

The age of the patients ranged from 0 to 57 years (median = 1 year), with a male–female ratio of nearly 3:1 (16 males vs. 6 females). Twenty patients (90.9%) were children (≤9 years old), including twelve (54.5%) infants (≤1 year old). The majority of the patients manifested leukocytosis (range 1–608 × 10^9/L), anemia (range 41–109 g/L) and low platelet counts (range 9–254 × 10^9/L). According to the high WBC index (23), nine (60.0%) of the fifteen cases with WBC count information were defined as high WBC levels. Twelve cases were not provided with a description of clinical features. Of the remaining 10 cases, five children (50.0%) had splenomegaly and hepatomegaly, and three patients (30.0%) had lymphadenopathy. Central nervous system involvement was observed in three children (30.0%), and skin involvement was observed in two cases (20.0%). All patients were diagnosed with AML (twenty children and two adults) according to the former FAB classification: five patients (three children and two adults, 22.7%) with M5, five children (22.7%) with M4, eight children (36.4%) with M2, one child (4.5%) with M1, and three children (13.6%) unknown. All the cases had available cytogenetic information, and chromosomal translocations (eleven cases) were the most common chromosomal rearrangements, followed by chromosomal insertions (nine cases). Among them, Xq24 (nine cases) and 11q23 (fourteen cases) were the most frequently involved chromosomal bands. Three cases (13.6%) demonstrated complex abnormalities. Eight patients (42.1%)
| Patient No. | Sex | Age | Hepatomegaly | Splenomegaly | CNS involvement | WBC/Hb/PLT (× 10^9/L) | Blood blasts (%) | Immunophenotype | Karyotype | Diagnosis | Treatment protocol | Treatment outcome | Survival (months) | Reference |
|------------|-----|-----|--------------|--------------|----------------|----------------------|-------------------|-----------------|-----------|-----------|-------------------|-----------------|-----------------|-----------|
| 1          | M   | 57  | No/No        | No           | No             | 12.3/79.0/105.0     | 69.0              | 92.0            | The leukemic cells expressed HLA-DR, CD117, CD33, CD13, CD38, CD15, CD64, and CD4. | 46, Y, t(X;11)[q24;q23] [4]/46,XY[6] | AML-M5 | IA regimen (resistance); then changed to decitabine combined with half-dose CAG regimen + high dose cytarabine | Resistance 2 months after diagnosis, CR 10 months after changing chemotherapy | 18 | Present |
| 2          | M   | 9   | Yes/Yes      | Yes          | Yes            | 3.0/93.0/245.0      | 2.0               | 27.5            | The leukemic cells expressed HLA-DR, CD33, CD38, CD15, CD64, and CD4. | 45,Y;del(X)(p21),del[11] t(X;11)[q24;q23],-20,add[2][q13][3] | AML-M2 | MAE regimen + intrathecal chemotherapy + BMT | CR 6 months after diagnosis | 43 | Present |
| 3          | M   | 16  | Yes/Yes      | No           | No             | 123.8/41.0/39.0     | 52.0              | 56.0            | The leukemic cells expressed HLA-DR, CD33, CD38, CD15, CD64, and CD4. | 46, Y, t(X;11)[q24;q23], del[7] [q21q31][13] /46,XY[2] | AML-M4 | HA regimen | Died 34 months after diagnosis | 36 | Present |
| 4          | M   | 0   | Yes/Yes      | No           | No             | 112.0/90.0/9.0      | 17.0              | 20.4            | The leukemic cells expressed HLA-DR, CD33, CD38, CD15, and CD64. | 46,X;ins(X;11)(p23;q24q12)[10] | AML-M5 | No chemotherapy | Died 0.5 month without chemotherapy | 0.5 | Present |
| 5          | F   | 3   | NA           | NA           | NA             | 163.2/NA/NA         | NA                | NA              | The leukemic cells expressed CD4, CD13, CD15, and CD33. | 46, XX, t(5,11)[q3:q23], add(X;q22) [12]/46, XX, t(5,11)[q13q23] [6]/47, XX, t(5,11), add(X;q22), +add(X) [1]/46, XX [1]. | AML-M2 | Chemotherapy + BMT | Died 9 months after diagnosis | 9 | (6) |

(Continued)
| Patient No. | Sex | Age (months) | Hepatomegaly / splenomegaly | CNS involvement | WBC/Hb/PLT (x 10⁹/L/g/L/ x 10⁹/L) | Blood blasts (%) | Bone marrow blasts (%) | Immunophenotype | Karyotype | Diagnosis | Treatment protocol | Treatment outcome | Survival months | Reference |
|------------|-----|--------------|-----------------------------|----------------|---------------------------------|-----------------|------------------------|-----------------|-----------|------------|---------------------|------------------|----------------|----------|
| 6          | M   | 7            | NA                          | NA             | 608.0/NA/NA                    | NA              | NA                     | The leukemic cells expressed CD4, CD13, and ins(X;11)(q22-24;q23) | 46, XY, del(11q) | AML-M2 | Chemotherapy + BMT | CR 35 months after diagnosis | 35 (6)         |
| 7          | F   | 6            | NA                          | NA             | 58.5/NA/NA                     | NA              | NA                     | The leukemic cells expressed CD4, CD13, CD33, and HLA-DR. | 46, X, add(X)(q23), AML-M1 del(11q) | AML-BFM 98 | CR 13 months after diagnosis | 13 (12)         |
| 8          | F   | 6            | NA                          | NA             | 280.0/NA/NA                    | NA              | NA                     | NA                           | 46,XX,ins(X;11)(q24;q23) | AML-M2 | Chemotherapy | Died during induction chemotherapy | NA (13) |
| 9          | F   | 20           | Yes/Yes                     | Yes            | 397.0/anemia/thrombocytopenia  | 91.0            | 46,XY,ins(X;11)(q22;q23)(t3,11) | The leukemic cells expressed CD33, CD15, CD11b, and HLA-DR. | 47,X, der(X);t(X;11) | AML-M4 | CR 1 month after diagnosis | Died during induction chemotherapy | 8 (14)         |
| 10         | M   | 10           | NA                          | NA             | 13.4/anemia/thrombocytopenia   | 13.0            | 46,Y(t(X;11)(q22;q23)(25))/46,XY[5] | The leukemic cells expressed CD11, CD13, CD15, CD33, and did not express lymphoid antigens. | 46,Y inv,ins(X;11)(q24;q33) | AML-M2 | CCG 2891 protocol + haploidentical transplant + BHAC regimen | PR 4 months after treatment, CR 7 years after diagnosis | 84 (13)         |
| 11         | M   | 29           | NA                          | NA             | 74.2                           | NA              | 46,Y inv,ins(X;11)(q24;q33) | The leukemic cells expressed CD13, CD33, HLA-DR, TdT, and CD7. | 46,Y inv,ins(X;11)(q24;q33) | AML-M5 | BHAC regimen | CR 1 month after diagnosis | 8 (14)         |
| 12         | M   | 8 months     | Yes/Yes                     | Yes            | 112.0/84.0/41.0                | 63.5            | 46,Y inv,ins(X;11)(q24;q33) | The leukemic cells expressed CD33, CD15, CD11b and CD36, and negative for CD34, CD13, CD14, CD7, CD19, and TdT. | 46,Y inv,ins(X;11)(q24;q33) | AML-M4 | 1-β-D-arabinofuranosylcytosine + daunomycin + etoposide + intrathecal chemotherapy | CR 1 month after diagnosis, died 5 months after relapse | 8 (15)         |
| No. | Patient Sex | Age (years) | Bone Involvement | CNS Involvement | WBC/Hb/PLT (×10^9/L) | Blood Blasts (%) | Bone marrow blasts (%) | Immunophenotype | Karyotype | Diagnosis | Treatment Protocol | Treatment Outcome | Survival (months) | Reference |
|-----|-------------|-------------|------------------|----------------|----------------------|-------------------|-------------------|----------------|-----------|-----------|------------------|----------------|----------------|-----------|
| 13  | M           | 3           | NA               | NA             | 2.4/3.5/4.6         | NA                | NA                | 46,XY,t(X;11)   | (q22;q23) | AML M2    | BMT               | CR               | 97 after diagnosis |
| 14  | M           | 12          | NA               | NA             | 2.2/2.5/3.6         | NA                | NA                | 46,XY,t(X;11)   | (q13;q23) | AML M4    | BMT               | Died             | 24.4 after diagnosis |
| 15  | M           | 12          | NA               | NA             | 2.2/2.5/3.6         | NA                | NA                | 46,XY,t(X;11)   | (q24;q23) | AML M5    | Chemotherapy      | CR               | 101.2 after diagnosis |
| 16  | M           | 6           | NA               | At diagnosis: 1.6/84.0/254.0 | At relapse: 9.5/115.0/85.0 | NA                | NA                | CD4 and CD13, did not express HLA-DR, CD33, CD11b, and other lymphocyte-associated antigens | 46,XY,t(X;11) (q24;q23) | AML M2    | BMT               | CR               | 61 months after diagnosis, died 11 months after relapse |
| 17  | F           | 12          | No/No            | No             | 1.0/93.0/81.0       | 28                | 82.0              | 47,X,add(X)(p11), +6,add[11](q23)[20].ish der(X)add(X)(p11)ins(X;11)(q24;q23), der(11)ins(11;?) (q22;?)ins(X;11) | (Continued) | AML M2    | ELAM 02 + BMT | CR                     | 96.9 after diagnosis |
| 18  | M           | 43          | No/No            | No             | 1.0/109.0/95.0      | NA                | NA                | CD33 and MPO. | 46,XY,t(X;11) | AML M5    | Palliative care only | CR               | 31 after diagnosis |

(Continued)
| Patient No. | Sex | Age (months) | Hepatomegaly / splenomegaly | CNS involvement | WBC/Hb/PLT (×10^9/L) | Bone marrow blasts (%) | Immunophenotype | Karyotype | Diagnosis | Treatment protocol | Treatment outcome | Survival months | Reference |
|------------|-----|--------------|----------------------------|-----------------|---------------------|------------------------|-----------------|-----------|-----------|-------------------|-----------------|-----------------|-----------|
| 19         | F   | 17           | NA                         | NA              | NA                  | NA                     | NA              | 47,X,add(X)(p11), +6,add(11)(q23)[20] | AML M2       | NA        | NA        | NA                  | 20              |
| 20         | M   | 12           | NA                         | NA              | NA                  | NA                     | NA              | 46,Yt(X;11)(q24;q23)[11]/46,XY[9] | AML          | NA        | NA        | NA                  | 20              |
| 21         | M   | 0            | NA                         | NA              | NA                  | NA                     | NA              | 46,Y,ins(X;11)(q24;13q23)[11] | AML          | NA        | NA        | NA                  | 20              |
| 22         | M   | 13           | No                         | 35.0/64.0/35.0  | 6.0                 | 73.0                   | The leukemic cells expressed CD45, CD33, CD15, CD34, CD64, HLA-DR, and MPO. In addition, a small population of blasts with monocytic differentiation (CD 641 and CD141) was detected. | 46,Y,ins (11;X)(q23; q24q22)[14]/46,del, i(10)[q10][6] | AML M4 | CCG 2891 protocol, Regimen C + intensive intrathecal chemotherapy + radiation to the extramedullary sites of disease + BMT | CR 2 months after diagnosis | 6              | (21)      |

M, male; F, female; CNS, central nervous system; WBC, white blood cells; Hb, hemoglobin; PLT, platelets; CAG, cytarabine + aclacinomycin + granulocyte colony stimulating factor (G-CSF); MAE, Mitoxantrone + cytarabine + etoposide; BMT, bone marrow transplantation; HA, homoharringtonine + cytarabine; BHAC, idarubicin + 1β-D-arabinofuranosylcytosine + thioguanine; CCG, the Children’s Cancer Group; AML-BFM, acute myeloid leukemia-Berlin-Frankfurt-Munster; CR, complete remission; PR, partial remission; NA, not available.
received chemotherapy alone. Nine patients (47.4%) received BMT. Eight patients (42.1%) died.

Of all 22 cases, 18 cases had clinical follow-up with a median period of 27.7 months (0.5–101.5 months). Kaplan-Meier survival analysis was performed on eighteen cases with complete follow-up information (Figure 4A). As of the final follow-up, the median survival time was 72 months.

To understand the impact of different clinical features on overall survival (OS), we grouped the patients according to the characteristics in Table 3, and groups with fewer than three patients and cases with incomplete follow-up information were not included in the statistics. The results showed that there was no statistical significance in OS between the infant group (≤1 year old) and pediatric group (>1 and <18 years old) (P = 0.4632, Figure 4B). The OS of patients with WBC levels ≥20.0 × 10^9/L was much shorter than that of patients with WBC levels < 20.0 × 10^9/L (P = 0.0072, Figure 4C). The patients with M4 had a shorter OS than those with M2 (P = 0.0212, Figure 4D). However, there were no significant differences in OS between the chromosomal translocation group and the chromosomal insertion group (P = 0.7095, Figure 4E) or between the patients who received chemotherapy alone and those who received BMT (P = 0.6958, Figure 4F).

Discussion

The KMT2A gene is a frequent target of rearrangement in human leukemia, especially in infant and pediatric leukemia (14, 24, 25). These rearrangements include fusions with many partner genes but rarely involve the gene SEPT6 on the X chromosome. The KMT2A gene and SEPT6 gene are vulnerable to damage to form translocations associated with infant AML.

In this study, we described four cases of AML with the KMT2A-SEPT6 fusion gene. The FAB subtypes were mainly M2, M4, and M5, which was consistent with the literature. To the best of our knowledge, only two adult patients have been reported, including one case in our series. Most of the cases that have been reported were children, and 54.5% were infant patients (≤1 year old). Among these cases, 60.0% had a high level of WBC, and 30% manifested central nervous system involvement, which were similar to the clinical features of KMT2A-rearranged AML patients. The findings of Balgobind et al. (26) showed that KMT2A-rearranged AML patients usually exhibit a high tumor burden, including organomegaly, a high median WBC count and central nervous system involvement. The present study included the largest number of KMT2A-SEPT6 cases to date. The patients’ NGS test results were not provided except ours, and we also tracked the patients’ molecular biological examination results. Three of four cases in our series had NRAS mutations, while one case with congenital AML did not.

NRAS G12V is required in the leukemia self-renewal process, independent of its effects on growth and survival (27). Compared with other subtypes of leukemia, acute leukemia with KMT2A translocations (such as KMT2A-AF4, and KMT2A-AF9) harbored the fewest number of mutations, in which NRAS mutations commonly co-occur (27, 28). In our series, we identified NRAS mutations in KMT2A-SEPT6-positive AML patients for the first time, and most of the mutation sites appeared at codons 12 and 145. The former site is a hotspot mutant of NRAS, and the latter site has also been reported (29, 30). The VAF of NARS mutation decreased as the patient’s condition improved, indicating that there was no underlying residual clonal hematopoiesis after chemotherapy. When the patients achieved CR, they also turned negative. The underlying connection between

TABLE 3  Clinical features of evaluable patients.

| Characteristic       | N (%) |
|----------------------|-------|
| Gender               |       |
| Male                 | 16 (72.7%) |
| Female               | 6 (27.3%)  |
| Age (years)          |       |
| ≤1                   | 12 (54.5%)  |
| >1 and <18           | 8 (36.4%)   |
| ≥18                  | 2 (9.1%)    |
| WBC count (×10^9/L)  |       |
| ≥20.0                | 9 (60.0%)   |
| <20.0                | 6 (40.0%)   |
| Symptom at presentation |   |
| Hepatosplenomegaly   | 5 (50.0%)  |
| CNS involvement      | 3 (30.0%)   |
| Lymphadenopathy      | 3 (30.0%)   |
| Skin involvement     | 2 (20.0%)   |
| FAB classification    |       |
| M1                   | 1 (4.5%)    |
| M2                   | 8 (36.4%)   |
| M4                   | 5 (22.7%)   |
| M5                   | 5 (22.7%)   |
| Unknown              | 3 (13.6%)   |
| Chromosomal abnormalities |   |
| Translocations       | 11 (50.0%)  |
| Insertions           | 9 (40.9%)   |
| Complex abnormalities | 3 (13.6%)   |
| Treatment protocol   |       |
| Chemotherapy alone   | 8 (42.1%)   |
| BMT                  | 9 (47.4%)   |
| No chemotherapy      | 2 (10.5%)   |
| Survival outcome     |       |
| Alive                | 11 (57.9%)  |
| Died                 | 8 (42.1%)   |

WBC, white blood cells; CNS, central nervous system; BMT, bone marrow transplantation.

Chen et al. /one.tnum/zero.tnum./three.tnum/three.tnum/eight.tnum/nine.tnum/fmed./two.tnum/zero.tnum/two.tnum/two.tnum./eight.tnum/nine.tnum/zero.tnum/nine.tnum/five.tnum/nine.tnum
NRAS mutations and KMT2A-SEPT6 and whether non-congenital KMT2A-SEPT6-positive AML patients all have NRAS mutations remain to be further studied in a larger cohort in the future.

KMT2A gene rearrangement in AML usually indicates poor prognosis (5, 31). The prognostic significance of NRAS mutations in AML patients remains unclear (32–35). Of the four cases in our series, one developed drug resistance at first, one suffered relapse after BMT and two died, showing unsatisfactory therapeutic effect. However, whether the outcomes of patients with KMT2A-SEPT6 were aggravated by the concurrence of NRAS mutations needs a follow-up study. Kaplan–Meier curves demonstrated that the pediatric group (>1 and <18 years old) did not show better OS than the infant group (≤1 year old). Age may not be an independent prognostic factor for survival. Most of the patients received chemotherapy, nine of them received BMT, but three of them eventually died. The OS of patients between the chemotherapy alone group and the BMT group did not show a significant difference, which suggested that BMT may not improve the survival time of such patients. This was consistent with several studies and meta-analyses that suggested that BMT does not improve survival in patients with KMT2A rearrangement (36, 37). In addition, chromosome rearrangement patterns had no significant effect on the OS of patients. However, we found that a higher white blood cell count at the initial diagnosis was associated with a shorter OS. Moreover, FAB classification also has an impact on the prognosis of patients. A trend for worse OS was observed in M4 patients.

Our study has some limitations. First, this was a retrospective study, coupled with a limited number of reported cases and incomplete clinical information, resulting in small sample sizes in some subgroups, which may lead to false negative results. Second, our NGS detection only covers the twenty most frequently mutated genes in AML, and the prognostic effects of some critical genes may be neglected. Third, gene mutation information in the reported cases was not available and cases in our series were detailed but limited by sample size. Therefore, our findings need to be combined with more cases for further analysis in the future.

In conclusion, the KMT2A-SEPT6 fusion gene was more commonly observed in pediatric patients diagnosed with AML. NRAS mutations were observed in these patients, most frequently of the NRAS G12V hotspot mutation. Whether NRAS mutations are related to the occurrence of KMT2A-SEPT6-positive AML is currently unclear. The prognosis was...
related to the subtype of leukemia and WBC levels, but may not be related to age. BMT may not improve survival in these patients. More cases should be accumulated and summarized to better understand the profile in KMT2A-SEPT6-positive AML.

Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

Ethics statement

The studies involving human participants were reviewed and approved by the Ethics Committee of Shengjing Hospital of China Medical University. Written informed consent to participate in this study was provided by the participants’ legal guardian/next of kin.

Author contributions

SF and FC performed the study concept and design. FC developed the methodology and wrote the paper. FC and YY acquired, analyzed and interpreted the data, and performed the statistical analysis. SF reviewed and revised the paper. All authors read and approved the final paper.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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