Acute Myeloid Leukemia in a Father and Son with a Germline Mutation of ASXL1

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Acute myeloid leukemia in a father and son with a germline mutation of ASXL1

Karen Seiter1*, Kyaw Htun1, Paul Baskind1 and Zach Liu2

Abstract

Background: Myelodysplastic syndromes and acute myeloid leukemia usually occur sporadically in older adults. More recently cases of familial acute myeloid leukemia and/or myelodysplastic syndrome have been reported.

Case presentation: Currently we report a father and son who both developed myelodysplastic syndrome that progressed to acute myeloid leukemia. Both patients were found to have the identical mutation of ASXL1 on nextgen sequencing of both hematologic and nonhematologic tissues.

Conclusions: These cases support the diagnosis of a germline mutation of ASXL1.

Keywords: Familial acute myeloid leukemia, Familial myelodysplastic syndrome, ASXL1, Nextgen sequencing

Background

Myelodysplastic syndromes and acute myeloid leukemia usually occur sporadically in older adults. These diseases also occur in younger patients with congenital syndromes such as Fanconi anemia, dyskeratosis congenita, and severe congenital neutropenia [1, 2]. More recently cases of familial acute myeloid leukemia (AML) and/or myelodysplastic syndrome (MDS) have been reported with mutations in RUNX1, CEBPA, GATA2, ANKRD26, SRP72, DDX41 or ETV6 [1, 2]. Currently we report a father and son who both developed MDS to AML. Both patients were found to have the identical mutation of ASXL1 on nextgen sequencing. The presence of the mutation in nonhematologic tissues supports the diagnosis of a germline mutation of ASXL1.

Case presentation

Son

A 46 year old man with no prior medical history was referred to us for evaluation of leucopenia and thrombocytopenia in June of 2012. The white blood count was 3200/mm³ with 51% neutrophils, 46% lymphocytes, 2% monocytes and 1% eosinophils. The hemoglobin was 13.1 g/dl and the platelet count was 103,000/mm³. The bone marrow was normocellular with 20% blasts and megalakaryocytic dysplasia. Flow cytometry demonstrated that the immunophenotype was CD13, CD33, CD11c, CD34, CD117, HLA-DR, CD71, CD41 (very dim), CD38, and CD9 (dim). Cytogenetics were normal, however only 4 metaphases were obtained. AML and MDS fluorescence in-situ hybridization (FISH) panels were normal. A diagnosis of high grade MDS evolving to AML was made.

The patient received induction therapy with high dose cytarabine, high dose mitoxantrone, and etoposide. Bone marrow evaluation upon count recovery showed a hypercellular marrow with 5% myeloblasts and dysplastic features, including pseudo-Pelger-Huet cells and dyserythropoiesis. The patient received a second cycle of chemotherapy with high dose cytarabine. The bone marrow then showed a normocellular marrow with maturing trilineage hematopoiesis, dyserythropoiesis and 1% blasts.

In September 2012 the patient underwent an allogeneic stem cell transplant using a fludarabine, melphalan, busulfan conditioning regimen and peripheral blood stem cells from his HLA identical sister. He has remained well, with full donor engraftment, for the past 3.5 years.

Father

In September 2016 the patient’s 75 year old father presented to us. He had been diagnosed with MDS elsewhere in January 2012. The initial bone marrow demonstrated
5% blasts and normal cytogenetics. He had received decitabine for 4.5 years. The patient was referred to us after his blood counts had worsened. The white blood count was 2900/mm³ with 56% neutrophils, 42% lymphocytes, 1% monocytes, and 1% eosinophils. The hemoglobin was 9.6 g/dl and the platelet count was 28,000/mm³. Bone marrow evaluation demonstrated AML with MDS-related changes, with 26% blasts and an immunophenotype of CD13, CD33, CD34, CD117, and HLA-DR. The patient was referred to us for decitabine for 4.5 years. The patient was referred to us for decitabine for 4.5 years. The patient was referred to us for decitabine for 4.5 years. The patient was referred to us for hospice care and expired shortly thereafter.

Nextgen sequencing was performed on the diagnostic bone marrow sample and a buccal swab from the father, and on peripheral blood, buccal swab, skin biopsy and archived bone marrow from the son. Genomic DNA was extracted and purified. Targeted sequencing was performed by using a combination of multiplexed PCR (AmpliSeq Hotspot primers) to generate libraries. Adapters were then ligated to the PCR products, where the sequences were tagged with specific barcodes. The barcoded libraries were then clonally amplified using emulsion PCR (emPCR). The emPCR was then purified using magnetic bead purification followed by semiconductor-based sequencing on an Ion Torrent PGM (Life Technology). The targeted gene panel developed in this laboratory includes the following 32 genes: ABL1, ASXL1, BRAF, CBL, CDKN2A, CEBPA, CREBBP, CSF1R, DNMT3A, ETV6, EZH2, FBXW7, FLT3, HRAS, IDH1, IDH2, JAK3, KIT, KRAS, NCOA2, NPM1, NRR2, PDGFR, PHF6, PTEN, RUNX1, SF3B1, SRSF2, TET2, TP53, U2AF1, and WT1. Each variant was analyzed manually using variant caller from Ion Torrent software (Life Technology) and cross-referenced with Ingenuity(tm) software (Qiagen) for bioinformatics. Evidence based categorization of the variants were performed by genomic analysis software (GenomOncology). This test is designed to detect alterations in genes that are clinically known to play a role in tumor genesis and provide prognostic value. For each gene, the minimum required coverage is 500 sequence reads based on bidirectional sequencing. The minimum acceptable frequency is 5%.

Both patients were enrolled on a clinical trial of familial leukemias that was approved by the Committee on the Protection of Human Subjects at New York Medical College. Both specifically consented to genetic testing. These included consent for publication.

Results of nextgen sequencing are summarized in Table 1. The father’s diagnostic bone marrow demonstrated a mutation in ASXL1 (c.2957A>G; p.N986S) with an allelic frequency of 50%. In addition a mutation in TP53 was detected with an allelic frequency of 12%. A buccal swab identified the same ASXL1 mutation at a frequency of 49.4%. We then tested the patient’s son for the same panel of mutations. His peripheral blood did not exhibit any mutations, however the blood was derived from his healthy sister (100% XX by FISH at the time of peripheral blood nextgen sequencing). A buccal swab on the son demonstrated the identical ASXL1 mutation (c.2957A>G; p.N986S) at a frequency of 21.7%. A skin biopsy demonstrated this mutation at a frequency of 41.6% and his archived first post chemotherapy bone marrow (with 5% blasts) demonstrated the mutation at a frequency of 50.5%.

**Table 1 Nextgen sequencing results**

| Patient | Date     | Source          | Gene  | Mutation               | Allelic burden |
|---------|----------|-----------------|-------|------------------------|----------------|
| Father  | 9/20/2016| Bone marrow     | TP53  | c.675delT; p.G226 fs   | 12.0%          |
|         |          |                 | ASXL1 | c.2957A>G; p.N986S    | 50.0%          |
| Son     | 10/26/2016| Buccal swab     | ASXL1 | c.2957A>G; p.N986S    | 49.4%          |
|         | 7/17/2012| Bone marrow (5% blasts) | ASXL1 | c.2957A>G; p.N986S    | 50.5%          |
|         | 11/4/2016| Buccal swab     | ASXL1 | c.2957A>G; p.N986S    | 21.7%          |
|         | 12/09/2016| Peripheral blood (100% donor) | Normal |                        |                |
|         | 1/13/2017| Skin biopsy     | ASXL1 | c.2957A>G; p.N986S    | 41.6%          |

Discussion and conclusions

Constitutive mutations of ASXL1 occur in the Bohring-Opitz syndrome, a rare condition characterized by facial anomalies, multiple malformations, severe intellectual disabilities and early death [3]. Somatic mutations of ASXL1 were first reported in patients with hematologic malignancies in 2009 [4]. Subsequent studies have since shown that mutations in ASXL1 occur in approximately 6% to 30% of patients with AML and in 15 to 20% of patients with MDS [5–7]. These mutations are more common in older patients and are more common in patients with secondary rather than de novo AML [8]. Mutations in ASXL1 have an adverse effect on survival in both MDS and AML [5, 9, 10].

In a mouse model developed by Abdel-Wahab, mice with germline complete deletion of ASXL1 (ASXL1−/−) were no longer viable by day 19.5 and exhibited microphthalmia/anophthalmia, cleft palates and multiple skeletal...
abnormalities [11]. Mice with hematopoietic –specific deletion of \textit{ASXL1} developed progressive leucopenia and anemia that was accompanied by an increase in erythroid precursor cells in both the bone marrow and spleen. Wang also reported that \textit{ASXL1} \textit{+/−} mice had developmental abnormalities including dwarfism, anophthalmia, and 80% embryonic lethality [12]. Surviving mice developed features of MDS. In this model, \textit{ASXL1} \textit{+/−} mice also developed an MDS-like phenotype indicating a haploinsufficient effect of \textit{ASXL1} in the pathogenesis of myeloid malignancies.

\textit{ASXL1} mutations have been reported in a limited number of patients with familial hematologic malignancies. Somatic mutation of \textit{ASXL1} was reported in a patient with chronic myelomonocytic leukemia in the setting of a germline mutation in \textit{ANKRD26} [13]. Somatic mutations of \textit{ASXL1} have also been reported in patients with germline mutations of \textit{GATA2} [14]. Hamadou described two sisters with NHL who were noted to have a presumed germline \textit{ASXL1} mutation in peripheral blood samples performed when they were in remission [15]. The sisters’ healthy mother and brother also carried this mutation.

Our two patients shared the identical \textit{ASXL1} mutation in multiple tissues. This specific mutation (c.2957A > G; p.N986S) has been reported in one case of Bohring-Opitz syndrome in the National Center for Biotechnology Information database. Its significance in hematologic malignancies to date is unclear; however it is a missense mutation in exon 12, a site of other known pathogenetic mutations in \textit{ASXL1} in hematologic malignancies [14]. Both patients were similar to the heterozygous mouse models in that their hematologic malignancies occurred after a prolonged latency period. The son’s earlier development of MDS/AML could be an example of anticipation, as is often seen in hereditary cancer syndromes. We believe this is the first reported case of a germline mutation of \textit{ASXL1} in association with MDS and AML.

Abbreviations

AML: Acute myeloid leukemia; CLAG: Cladribine, cytarabine, filgrastim; DNA: Deoxyribonucleic acid; FISH: Fluorescence in-situ hybridization (FISH); HLA: Human leukocyte antigen; MDS: Myelodysplastic syndrome; PCR: Polymerase chain reaction

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Availability of data and materials

Please contact author for data requests.

Authors’ contributions

KS: designed the study, collected and analyzed the data and wrote the manuscript; KH: collected and analyzed the data; PB: collected and analyzed the data; ZL: collected and analyzed the data and wrote the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The study was approved by the Institutional Review Board of New York Medical College. The patients consented to participate.

Consent for publication

The patients consented to publication.

Competing interests

The authors declare that they have no competing interests.

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References

1. Feurstein S, Drazer MW, Godley LA. Genetic predisposition to leukemia and other hematologic malignancies. Semin Oncol. 2016;43:598.
2. Chrupek JE, Pytel K, Kanchi KL, Shao J, Koboldt D, Miller CA, et al. Genomic analysis of germ line and somatic variants in familial myelodysplasia/acute myeloid leukemia. Blood. 2015;126:2484.
3. DangiofbloSB, Wilson A, Jobanputra Anyanyeboa K, Bohring-Opitz syndromewith a new \textit{ASXL1} pathogenic variant: review of the most prevalent molecular and phenotypic features of the syndrome. Am J Med Genet. 2015;167A:3161.
4. Gelsi-Boyer V, Trouplin V, Adelade J, Bonanseaa J, Cervera N, Carbucia N, et al. Mutations of polycomb-associated gene \textit{ASXL1} in myelodysplastic syndromes and chronic myelomonocytic Leukaemia. Br J Haematol. 2009;145:788.
5. Bejar R, Stevenson K, Abdel-Wahab O, Gallin N, Nilsson B, Garcia-Manero G, et al. Clinical effect of mutations in myelodysplastic syndromes. N Engl J Med. 2011;364:2496.
6. Devillier R, Mansat-De Mas V, Gelsi-Boyer V, Demur C, Murati A, Corre J, et al. Role of \textit{ASXL1} and \textit{TP53} mutations in the molecular classification and prognosis of acute myeloid leukemias with myelodysplasia-related changes. Oncotarget. 2015;6:8388.
7. Haferlach T, Nagata Y, Grossmann V, Okuno Y, Bacher U, Nagae G, et al. Landscape of genetic lesions in 944 patients with myelodysplastic syndromes. Blood. 2014;126:2141.
8. Nazha A, Zarzour A, Al-Issa K, Radioyevich T, Carraway HE, Hirsch CM, et al. The complexity of interpreting genomic data in patients with acute myeloid leukemia. Blood Cancer. 2016;6:e10.
9. Metzeler KH, Becker H, Maharry K, Radmacher MD, Kohlschmidt J, Mrozek K, et al. \textit{ASXL1} mutations identify a high-risk subgroup of older patients with primary cytogenetically normal AML within the EML favorable genetic category. Blood. 2011;118:6920.
10. Thol F, Friesen I, Damm F, Yun H, Weissinger EM, Krauter J, et al. Prognostic significance of \textit{ASXL1} mutations in patients with myelodysplastic syndromes. J Clin Oncol. 2011;29:2499.
11. Abdel-Wahab O, Gao J, Adli M, Dey A, Trimarchi T, Chung YR, et al. Deletion of \textit{ASXL1} results in myelodysplasia and severe developmental defects in vivo. J Exp Med. 2013;210:2641.
12. Wang J, Li Z, He Y, Pan F, Chen S, Rhodes S, et al. Loss of \textit{ASXL1} leads to myelodysplastic syndrome-like disease in mice. Blood. 2014;123:541.
13. Perez Botero J, Oliveira J, Chen D, Reichard KK, Wiswanatha DS, Nguyen PL, et al. \textit{ASXL1} mutated chronic myelomonocytic leukemia in a patient with familial thrombocytopenia secondary to germline mutation in \textit{AWD26}. Blood Cancer J. 2015;5:e315.
14. West RR, Hsu AP, Holland SM, Cuellar-Rodriguez J, H Hickstein DD. Acquired \textit{ASXL1} mutations are common in patients with inherited \textit{GATA2} mutations and correlate with myeloid transformation. Haematologica. 2014;99:276.
15. Hamadou WS, Abed RE, Besbes S, Boudoun Y, Fabre A, Youssef YB, et al. Familial hematological malignancies: \textit{ASXL1} gene investigation. Clin Transl Oncol. 2016;18:385.