Mutation profile of acute myeloid leukaemia in a Chinese cohort by targeted next-generation sequencing

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Abstract

Background: Acute myeloid leukaemia (AML) results from the clonal expansion of blast cells of myeloid origin driven by genomic defects. The advances in next-generation sequencing (NGS) have allowed the identification of many mutated genes important in the pathogenesis of AML.

Aims: In this study, we aimed to assess the mutation types and frequency in a Chinese cohort presenting with de novo AML cohort using a targeted NGS strategy.

Methods: In total, we studied samples from 87 adult patients with de novo AML who had no prior history of cytotoxic chemotherapy. Samples were evaluated using a 120-gene targeted NGS panel to assess the mutation profile.

Results: Of the 87 AML patients, there were 60 (69%) with a normal karyotype. 89.7% of patients had variants, with an average of 1.9 mutations per patient (range: 0–5 mutations per patient). DNMT3A variants were the most common, being detected in 33 patients (37.9%). NPM1 (34.5%), IDH1/2 (24.1%) and FLT3-ITD (20.7%) mutations was the next most common. Of the patients with DNMT3A mutations, 24.2% also had mutations NPM1 and FLT3-ITD and 6.1% NPM1, FLT3-ITD and IDH mutations.

Conclusion: Both DNMT3A and NPM1 mutations were more common than in other Chinese and Western AML cohorts that have been studied. DNMT3A mutations tended to co-occur with NPM1 and FLT3-ITD mutations and were most commonly seen with a normal karyotype.

KEYWORDS
Chinese AML, DNMT3A, mutation frequency, next-generation sequencing

1 INTRODUCTION

Acute myeloid leukaemia (AML) is a group of heterogeneous diseases characterised by distinct clinical, morphological, cytogenetic and genetic features. AML is classified as a unique disease entity in the current 2017 World Health Organization (WHO) classification of tumours of haematopoietic and lymphoid tissues1,2,3 based on specific recurring genetic abnormalities for predicting prognosis and treatment response. Specific mutations that are included in the classification are NPM1 and CEBPA. Recurrent chromosomal structural abnormalities are now confirmed as diagnostic and prognostic markers, which indicate that acquired genetic abnormalities are essential in leukaemogenesis.4,5,6 However, many cytogenetically normal AML (NK-AML) cases that lack recurrent structural abnormalities, and account for 40% to
50% of de novo adult AML are associated with an intermediate clinical outcome. It is known that the prognosis of AML patients is significantly affected by multiple genetic mutations and many leukaemias are composed of multiple subclones with different responses to treatment protocol. In the Cancer Genome Atlas (TCGA) group, 23 significantly mutated genes were identified by whole-genome or whole-exome sequencing in 200 AML patients. These include genes that are well known to be related in AML pathogenesis (e.g., DNMT3A, FLT3, NPM1, IDH1, IDH2 and CEBPA). For AML risk classification, it is important to study the mutations in multiple genes simultaneously because of complicated interactions with different pathways in leukaemogenesis. The majority of data regarding the frequencies of specific AML mutations have been derived from Caucasian populations. These have shown frequencies for the most common variants ranging from 26.3% to 33.0% for FLT3, 23.0% to 28.0% for NPM1 and 22.1% to 26.0% for DNMT3A. Other populations groups that have
| Patient gender/age (years) | WHO/FAB diagnosis | Karyotype                  | DNMT3A mutations          | NPM1 mutations | FLT3-ITD mutations | IDH1/2 mutations |
|---------------------------|-------------------|---------------------------|---------------------------|----------------|-------------------|-----------------|
| F/54                      | AML without maturation/AML M1 | 46, XX[18] | p.Arg882Cy (R882C) | Negative | Positive (57 base pairs duplication) | IDH1: p.Arg132His (R132H) |
| M/48                      | Acute myelomonocytic leukaemia/AML M4 | 46, XY[22] | p.Pro904Leu (P904L) | Negative | Negative | Negative |
| M/78                      | AML with maturation/AML M2 | 46, XY[19] | p.Arg882His (R882H) | Negative | Positive (8 base pairs duplication) | Negative |
| F/54                      | AML with maturation/AML M2 | 47, XX,+?15[2]/46, XX[18] | p.Arg882His (R882H) | Negative | Negative | IDH2: p.Arg140Gln (R140Q) |
| F/46                      | AML without maturation/AML M1 | 46, XX[20] | p.Ser714Cys (S714C) | Negative | Negative | IDH2: p.Arg172Lys (R172K) |
| F/70                      | Acute myelomonocytic leukaemia/AML M4 | 46, XX[19] | p.Lys299Asnfs (K299N) | Negative | Negative | Negative |
| F/53                      | AML without maturation/AML M1 | 47, XX, t(8;21)[q22; q22]+15[19] | p.Lys299Asnfs (K299N) | Negative | Positive (39 base pairs duplication) | Negative |
| F/72                      | Acute monocytic leukaemia/AML M5b | 46, XX[20] | p.Arg882 Pro (R882P) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Positive (18 base pairs duplication) | Negative |
| F/43                      | Acute myelomonocytic leukaemia/AML M4 | 46, XX[20] | p.Arg882 His (R882H) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Positive (48 base pairs duplication) | Negative |
| M/52                      | AML without maturation/AML M1 | 46, XY, t(12;12)[q23; q24.1][20] | p.Cys710Tyr (C710Y) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Positive (42 base pairs duplication) | Negative |
| M/66                      | AML without maturation/AML M1 | 46, XY[18] | p.Glu545* | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Positive (87 base pairs duplication) | IDH2: p.Arg140Gln (R140Q) |
| M/45                      | Acute myelomonocytic leukaemia/AML M4 | 46, XX[20] | p.Arg882Cys (R882C) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | IDH2: p.Arg140Gln (R140Q) |
| F/62                      | AML without maturation/AML M1 | 46, XX[21] | p.Arg882His (R882H) | Negative | Negative | IDH2: p.Arg172Lys (R172K) |
| M/59                      | AML with maturation/AML M2 | 46, XY,t(10;11)[q23; p15][1]/46, idem, add(6)[p25][19] | p.Arg882His (R882H), p.Lys299Asnfs (K299N) | Negative | Positive (27 base pairs duplication) | Negative |
| F/70                      | AML with maturation/AML M2 | 46, XX[20] | p.Ser714Cys (S714C), p.Glu545* | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Positive | IDH1: p.Arg132His (R132H) |
| F/78                      | AML without maturation/AML M1 | 46, XX[20] | p.Arg882His (R882H), p.Glu545* | Negative | Negative | Negative |
| Patient gender/age (years) | WHO/FAB diagnosis | Karyotype | DNMT3A mutations          | NPM1 mutations | FLT3-ITD mutations | IDH1/2 mutations |
|---------------------------|-------------------|-----------|---------------------------|----------------|---------------------|------------------|
| M/65                      | Acute myelomonocytic leukaemia/AML M4 | 46, XY[20] | P.Glu545*                 | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | IDH1: p.Arg132His (R132H) |
| M/72                      | AML with maturation/AML M2 | 46, XY[20] | p.Trp893Val, p.Glu545*    | Negative        | Negative            | Negative         |
| F/87                      | AML with myelodysplasia-related changes (multilineage dysplasia) | 46, XX[20] | p.Glu545*                 | Negative        | Negative            | Negative         |
| M/50                      | AML without maturation/AML M1 | 46, XY[20] | p.Arg882His (R882H), p.Glu545*, p.Trp893Valfs | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | Negative         |
| M/80                      | AML without maturation/AML M1 | 46, XY[20] | p.Cys911Tyr (C911Y), p.Glu545* | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | Negative         |
| F/49                      | AML without maturation/AML M1 | 46, XX[20] | p.Glu545*                 | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Not done | Negative         |
| F/82                      | AML with myelodysplasia-related changes (multilineage dysplasia) | 46, XX[20] | p.Glu545*                 | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | Negative         |
| M/81                      | AML with maturation/AML M2 | 46, XY[20] | p.Glu545*                 | Negative        | Negative            | Negative         |
| M/88                      | Acute myelomonocytic leukaemia/AML M4 | 46, XY[20] | p.Ser638Cys (S638C) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Positive | Negative         |
| M/50                      | AML without maturation/AML M1 | 46, XY[20] | pThr503Asnfs (T503N) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | Negative         |
| M/41                      | AML without maturation/AML M1 | 46, XY[20] | p.Arg882His (R882H) | p.Trp288Cysfs*12 (W288C) (mutation A with TCTG insertion) | Negative | Negative         |
| F/42                      | AML with maturation/AML M2 | 46, XX[20] | p.Arg885Trp (R885W), p.Val649Met | Negative        | Negative            | IDH1: p.Arg132Cys (R132C) |
been studied are Japanese, Korean and Singaporean. The genes that are involved in AML in these populations are the same, but with some differences in frequency. Few studies have assessed AML mutations in Chinese populations. These reports have shown the mutational frequencies ranging from 12.5% to 14.0% for DNMT3A and 15.0 to 15.9 for NPM1. In the present study, we have progressed this by studying an adult Chinese population presenting with de novo AML using a next-generation sequencing (NGS) platform. This targeted approach assessed the mutation profile and frequency of 120 genes associated with myeloid malignancies. The purpose of this study was to assess the frequency of mutations in the genes most commonly associated with aberrations in myeloid neoplasms in a population of Chinese patients with de novo AML.

2 | MATERIALS AND METHODS

2.1 | Patients and samples

In total 87 adult Chinese patients with de novo AML without any prior history of cytotoxic chemotherapy were studied. These patients were diagnosed with AML between 2004 and 2014 in two acute hospitals in Hong Kong and classified according to the 2001 and 2008 WHO classification. Diagnoses were made on bone marrow morphology and immunophenotyping, using standard methods.

Cytogenetic analysis was performed on short-term unstimulated synchronised culture using fluorodeoxyuridine (FdU) of the BM or PB samples. Karyotypes were analyzed after Giema-banding and reported in the International System for Human Cytogenetic Nomenclature. Patients were divided into favourable, intermediate and adverse risk groups according to their cytogenetic results and also the incorporation of molecular analyses as recommended by European LeukaemiaNet (ELN) in 2010.

DNA extraction was performed on the buffy coat of PB or BM samples using AllPrep DNA/RNA Mini Kit (Qiagen, Valencia, California) according to the manufacturer's protocol. FLT3-ITD mutations were analysed by conventional PCR based molecular method. FLT3-ITD mutations in exons 14 and exon 15 were detected by PCR and ITD was then confirmed by sequencing as previously described. NPM1 exon-12 mutations were detected by PCR and then fragment analysis was performed by ABI 3130 genetic analyzer (Applied Biosystems). The results were analyzed with GeneMapper Software Version 4.0 (Applied Biosystems) as previously described.

2.2 | Massively parallel sequencing

Ion AmpliSeq Designer was used to create an Ion AmpliSeq Custom Panel (Thermo Fisher Scientific). Twenty nanograms of DNA was used to create the amplicon library for sequencing the whole exons of 120 genes (Appendix S1) that are involved with myeloid disorders as previously described. The completed library was prepared using an Ion AmpliSeq Library Kit 2.0 and the custom primer panel (Thermo
Fisher Scientific) following the manufacturer’s instruction. Barcoded libraries were measured using the Qubit dsDNA HS Assay Kit and the Qubit 2.0 Fluorometer (Thermo Fisher Scientific). The barcoded libraries were clonally amplified by emulsion PCR (E-PCR) on Ion Sphere Particles (ISPs) using an Ion PI Template OT2 200 Kit v2 and the Ion OneTouch 2 System (Thermo Fisher Scientific). Enrichment of the template-positive ISPs was achieved by the Ion OneTouch 2 ES (Thermo Fisher Scientific). Sequencing of the enriched ISPs was prepared on an Ion Proton Sequencer (Thermo Fisher Scientific) using Ion Proton I (PI) chip and Ion PI Sequencing 200 Kit (Thermo Fisher Scientific) as previously described. Torrent Suite Software Version 4.0 (Thermo Fisher Scientific) was used for signal processing, base calling, sequence alignment, variant calling and to generate run metrics. Variants were confirmed with Variant Caller Version 4.0 (Thermo Fisher Scientific) and align reads to the reference human genome build 19. Ion Reporter Version 4.0 (Thermo Fisher Scientific) was used to annotate the variants and detect amino acid changes and diagnostic significance.

The following filtering criteria were used for the NGS variant mutations: Intron and exonic synonymous variants were removed while exonic non-synonymous variants were retained for analysis. For polymorphic variants, a minor allelic frequency ≥ 1% and/or found in single nucleotide polymorphism (SNP) Database were excluded. In addition, a variant allele frequency (VAF) > 5% was used as the cutoff.

2.3 | Statistical analysis

Chi-square test was used to analyze the association between different gene mutations and also for other categorical variables. Two-sided p value < .05 was considered statistically significant. All statistical analyses were performed with SPSS Version 26.0 (SPSS Inc, Chicago, Illinois).

3 | RESULTS

3.1 | Clinical and cytogenetic characteristics

There were 45 males and 42 females (M:F ratio = 1.07:1) newly presenting adult Chinese patients with de novo AML were included in the study. The mean age was 62 years (range: 20–91 years) with 39 patients (45%) being under the age of 60 years. Cytogenetic results showed 60 patients (60/87, 69.0%) with a normal karyotype (NK-AML) and 27 patients (27/87, 31.0%) were abnormal (AK-AML).

| TABLE 2 Co-occurrence of gene mutations in acute myeloid leukaemia (AML) patients |
|---------------------------------|-------------|-------------|-------------|-------------|
|                                | NPM1        | FLT3-ITD    |              |              |
|                                | WT          | Mutant      | p            | WT          | Mutant      | p            |
| DNMT3A                         | WT          | 41          | 13           | .009        | DNMT3A      | WT          | 48           | 6            | .002         |
|                                | Mutant      | 16          | 17           |             | Mutant      | 18          | 12           |              |             |
| IDH1/2                         | WT          | 43          | 11           | .293        | IDH1        | WT          | 50           | 4            | .460         |
|                                | Mutant      | 23          | 10           |             | Mutant      | 29          | 4            |              |             |
| IDH2                           | WT          | 47          | 7            | .508        | DNMT3A      | WT          | 47           | 7            | .908         |
|                                | Mutant      | 27          | 6            |             | Mutant      | 29          | 4            |              |             |
| NPM1                           | WT          | 50          | 19           | .008        | IDH1/2      | WT          | 45           | 21           | .354         |
|                                | Mutant      | 7           | 11           |             | Mutant      | 12          | 9            |              |             |
| IDH1                           | WT          | 53          | 26           | .333        | IDH2        | WT          | 49           | 25           | .743         |
|                                | Mutant      | 4           | 4            |             | Mutant      | 8           | 5            |              |             |
| IDH2                           | WT          | 66          | 8            | .213        |             |             |              |              |              |
|                                | Mutant      | 13          | 0            |             |             |              |              |              |              |

Abbreviation: WT, wild type.
3.2 Mutation profile

After applying the filtering criteria, an average of 1.9 mutations were detected per patient (range: 0–5 mutations) and variants were seen in 78/87 (89.7%) of all patients (Figure 1). Of these 60/87 (69%) had a normal karyotype. Because of the lack of matched constitutional normal control material, we could not confirm that all gene mutations were somatic mutations because some may be rare SNPs. DNMT3A mutations were detected in 33/87 (37.9%) of patients (mean age 61 years). The other commonly mutated genes were NPM1 (30/87, 34.5%), IDH1/2 (21/87, 24.1%) and FLT3-ITD (18/87, 20.7%) mutations (Figures 2, 3 and Table S1). Of the 33 patients with DNMT3A mutations (Table 1), 29 patients (29/33, 87.9%) had a normal karyotype. We also found that DNMT3A mutations had the highest frequency in NK-AML patients (33/60, 55%). The most frequent mutations were located in R882, with R882H mutation being the most common (9/13, 69.2%), followed by R882C mutation (3/13, 23.1%). DNMT3A mutations tended to co-exist with other mutations (Table 2). Among the patients with DNMT3A mutations, six patients had concomitant NPM1 and FLT3-ITD mutations. In addition, two patients had IDH1/2 mutation that co-occur with NPM1 and FLT3-ITD mutations.

The NPM1 mutations detected by NGS (30/87 or 34.5% patients) were successfully validated by conventional molecular methods. The majority of NPM1 mutations (26/30, 86.7%) were type A mutations (insertion of TCTG).27 There were two patients (2/30, 6.7%) with type D (insertion of CCTG), one (1/30, 3.3%) was type B (insertion of CATG) and another (1/30, 3.3%) with the type DD-3 (insertion of CAGA) mutations. Of these 30 patients NPM1-mutated cases, 29 (29/30, 96.7%) had a normal karyotype. NPM1 mutations were frequently concurrent with FLT3-ITD mutations (p = .008) (Table 2). There were FLT3-ITD mutations found in 18 of the 30 patients (18/87, 20.7% of all), of which 14 patients had a normal karyotype.

IDH1 mutations were detected in 8 patients (8/87, 9.2%) and all cases had a normal karyotype type-AML (NK-AML). All IDH1 mutations were located in R132 with R132H mutations being the most common (5/8, 62.5%) and followed by R132C mutations (3/8, 37.5%). IDH2 mutations were detected in 13 patients (13/87, 14.9%), 11 (84.6%) of whom had a normal karyotype. In 10 patients (10/13, 76.9%) IDH2 mutations were located in R140 with all of being R140Q mutations. IDH1 and IDH2 mutations did not co-exist in our cohort (Table 2).

4 DISCUSSION

In this study, we used a targeted NGS panel to sequence the whole exons of 120 genes known to be mutated in myeloid neoplasms, in 87 newly diagnosed Chinese adult patients with AML. Mutations were detected in 78/87 (89.7%) patients with an average of 1.9 mutations per patient. Of note, was that 60/87 (69%) patients with variants had NK-AML. The most frequently mutated genes were similar to those previously reported Caucasian AML cohorts, including DNMT3A, NPM1, IDH1/2 and FLT3-ITD.6,11,12 We showed higher gene mutation frequencies than reported for other Chinese cohorts.16,17,18 This was seen for both DNMT3A (37.9% vs. 12.5%–14.0%) and NPM1 (34.5% vs. 15.0%–15.9%) mutations.

We found DNMT3A was the most common mutated gene, present in 37.9% of patients, almost double the frequencies in other reports.6,11,12 Other Chinese studies16,17,18 had shown DNMT3A mutation frequencies of 12.5%–14.0% which were slightly less than the Caucasian populations and other Asian population countries (Korea: 17.5%; Japan: 19.0%; Singapore: 21.0%).13,14,15 Although the frequency of DNMT3A mutations was high, a single nucleotide change in R882 was commonly seen (39.4%), from arginine to histidine (R882H mutation, 69.2%) or cysteine (R882C, 23.1%), as reported in other studies.12,13,28 DNMT3A mutations seemed to co-occur with NPM1 (p = .009) and FLT3-ITD mutations (p = .002). Among the AML patients with DNMT3A mutations, 6.1% of patients had concomitant NPM1, FLT3-ITD and IDH mutations while 24.2% AML patients had triple mutations of DNMT3A, NPM1 and FLT3-ITD. These findings were in line with published reports on DNMT3A in patients with AML.6,11,13,29,30 The high frequencies of DNMT3A and NPM1 mutations in our study may be due to the high incidence of normal karyotype (69.0%). However, those with DNMT3A mutations were not in the older age group, as the mean age of this cohort was 61 years. Further those with concurrent DNMT3A and NPM1 mutations (n = 17) also were not of older age (mean = 60.2 years). These findings therefore do not support those of previous reports.6,13,30 that show that patients with DNMT3A mutations were generally older in age. A larger cohort is needed to further investigate these relationships and explore the different gene mutation frequencies are related to genetic background, lifestyle, environment and other factors.

DNMT3A is an epigenetic regulator catalyses DNA methylation in CpG islands and regulates the gene silencing processes.31 It is vital in normal haematopoietic stem cell differentiation32 and self-renewal and its mutation produces a sufficient amount of preleukaemic stem cells which finally convert into AML33. Mutations in DNMT3A and IDH1/2, genes that encode epigenetic modifiers, are present in the early pre-leukaemic cells and these “founder” mutations can be implicated as functional components of AML evolution. These genes are frequently mutated in elderly patients with clonal haematopoiesis.11 NPM1 mutations are regarded as secondary events and usually occur after DNMT3A and IDH1 mutations. These suggest that development of AML will follow specific and ordered evolutionary processes. Recent studies showed that DNMT3A R882 mutation exerts a dominant-negative effect. The mutant protein then interferes with the remaining normal DNMT3A to form active tetramers that reduces the enzyme activity and hypomethylation at specific cytosine-guanine dinucleotides in early AML cells.34

NPM1 is a nuclear protein involving ribosome biogenesis, DNA repair and prevent apoptosis.23 The frequency of NPM1 mutations was higher than in other Chinese cohorts (15.0%–15.9%).16,17,18 We found NPM1 mutations in 34.5% of our patients with the majority (86.7%) being type A mutations (insertion of TCTG). Of these, 96.7% had a normal karyotype type-AML.

The combined frequency of IDH1/2 mutations was 24.1%, in line with other publications.6,11 IDH1 mutations were found in 9.2% of the
AML patients and all cases had a normal karyotype (NK-AML). It has been reported that patients with IDH1 mutations are commonly older. This was not the case in the present study where the mean age of patients with IDH1/2 mutations was 60 years (range 42–85 years), consistent with the full cohort with mean age of 62 years. All IDH1 mutations detected were single nucleotide change in R132 with IDH2 mutations being the most common (62.5%). Deng et al. showed that R132 IDH1 mutations cause the encoded enzyme to acquire the novel ability to convert alpha-ketoglutarate (α-KG) to 2-hydroxy-glutarate (2HG). The increased cellular 2HG levels will cause inhibition of α-KG-dependent enzymes that are important for the demethylation of DNA. IDH2 mutations were more common, detected in 14.9% of the AML patients with 84.6% having normal karyotype. The most common IDH2 mutation caused changes of R140 with all cases were R140Q mutation. No patients had both IDH1 and IDH2 mutations, in keeping with previous reports, that these mutations are mutually exclusive. Studies showed that IDH1 and IDH2 seem to act as an epigenetic role in histone and DNA methylation. IDH1 and IDH2 mutations then cause a hypermethylation phenotype in leukaemia and inhibit haematopoietic stem cell differentiation.

Several studies have shown that AML patients with DNMT3A mutations had poorer clinical outcomes compared with the wild-type DNMT3A. DNMT3A mutations were also associated with worse survival for AML patients with a normal cytogenetic and those with an intermediate-risk profile. For the unique subgroup of AML patients with concomitant DNMT3A, NPM1 and FLT3-ITD mutations, they had the poorest prognosis. Since DNMT3A, NPM1 and FLT3 mutations belong to the three separate classes of mutations, this suggests the possible interaction between different classes of gene mutation in AML pathogenesis.

For our study, we only performed large-scale gene sequencing on the leukaemia samples. Although the use of matched normal specimens may be important in the identification of recurrent variants, it is not necessary when a good filtering system has been established. We used “population frequency” approach which is the percentage of the samples with mutation in the database of sample sequenced in order to filter out common benign variants. This is important in filtering out the common germline polymorphisms and homopolymer-related artefacts with particular high frequency.

In conclusion, simultaneous screening of multiple gene mutations using a 120-gene targeted NGS approach has identified high frequencies of genomic variants in adult de novo AML. In addition, we have shown that DNMT3A is the most common mutation in Chinese AML patients. The frequency of both DNMT3A and NPM1 mutations are higher than in other published studies of Chinese patients, three and twofold, respectively. DNMT3A mutations tended to co-occur with NPM1 and FLT3-ITD mutations in patients with NK-AML.

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CONFLICT OF INTEREST
The authors have stated explicitly that there are no conflicts of interest in connection with this article.

AUTHOR CONTRIBUTIONS
B.M.W.L.: Conceptualization; methodology; investigation; formal analysis; resources; data curation; writing-original draft; writing-review and editing. Y.L.K.: Conceptualization; methodology; formal analysis; resources; writing-original draft. W.N.E.: Conceptualization; methodology; validation; formal analysis; resources; supervision; project administration; funding acquisition; writing-original draft; writing-review and editing. B.B.G.: formal analysis. J.A.J.M.: formal analysis, manuscript drafting.

ETHICS STATEMENT
The study protocol was approved by the Research Ethics Committee, Hospital Authority, Hong Kong. The Ethics Committee had waived the requirement for informed consent because archival buffy coat samples were used and all data had been fully anonymized

DATA AVAILABILITY STATEMENT
The data that support the findings of this study are available from the corresponding author upon reasonable request.

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