



Original article

Identifying the similarities and differences between single nucleotide polymorphism array (SNPa) analysis and karyotyping in acute myeloid leukemia and myelodysplastic syndromes



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ABSTRACT

Objective: To standardize the single nucleotide polymorphism array (SNPa) method in acute myeloid leukemia/myelodysplastic syndromes, and to identify the similarities and differences between the results of this method and karyotyping.

Methods: Twenty-two patients diagnosed with acute myeloid leukemia and three with myelodysplastic syndromes were studied. The G-banding karyotyping and single nucleotide polymorphism array analysis (CytoScan[®] HD) were performed using cells from bone marrow, DNA extracted from mononuclear cells from bone marrow and buccal cells (BC).

Results: The mean age of the patients studied was 54 years old, and the median age was 55 years (range: 28–93). Twelve (48%) were male and 13 (52%) female. Ten patients showed abnormal karyotypes (40.0%), 11 normal (44.0%) and four had no mitosis (16.0%). Regarding the results of bone marrow single nucleotide polymorphism array analysis: 17 were abnormal (68.0%) and eight were normal (32.0%). Comparing the two methods, karyotyping identified a total of 17 alterations (8 deletions/losses, 7 trisomies/gains, and 2 translocations) and single nucleotide polymorphism array analysis identified a total of 42 alterations (17 losses, 16 gains and 9 copy-neutral loss of heterozygosity).

Conclusion: It is possible to standardize single nucleotide polymorphism array analysis in acute myeloid leukemia/myelodysplastic syndromes and compare the results with the abnormalities detected by karyotyping. Single nucleotide polymorphism array analysis increased the detection rate of abnormalities compared to karyotyping and also identified a new set of abnormalities that deserve further investigation in future studies.

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Introduction

Acute myeloid leukemia (AML) is a group of clonal disorders resulting from acquired somatic genetic lesions accumulated in hematopoietic progenitors in a step-wise fashion throughout life. The mutations give rise to a malignant hematopoietic clone; malignant hematopoiesis subsequently suppresses normal hematopoiesis.^{1,2}

The myelodysplastic syndromes (MDS) are a group of clonal hematopoietic stem cell disorders involving cytogenetic changes, and gene mutations that occur in a multistep process, with widespread gene hypermethylation at advanced stages. MDS are characterized by ineffective hematopoiesis leading to blood cytopenias or, in some cases, evolution to AML.³

The detection of chromosomal abnormalities in AML/MDS supports the diagnosis, classification, prognostic stratification, therapy option, treatment monitoring and better understanding of the disease's biology.^{3,4} The consequence of chromosomal investigations is major advances in the treatment and survival of patients. However, about 50% of AML/MDS have either normal karyotypes or non-recurrent chromosomal abnormalities, which prevents a better characterization of the disease.^{5,6} Therefore, an increased abnormality detection rate by other methods and improved understanding of pathogenesis are urgently needed.^{1,7,8}

In this context, the single nucleotide polymorphism array (SNPa) method, also referred to as molecular karyotyping, is a sensitive technology used to perform high-resolution genome-wide DNA copy number analysis and to detect segmental regions of homozygosity, known as regions of copy-neutral loss of heterozygosity (CN-LOH).^{2,9}

SNPa analysis uses millions of markers (probes comprise 25-mer oligonucleotides) across the entire human genome.¹⁰ There are two types of probes: non-polymorphic probes, used only for assessing copy number variations (CNV) (Figure 1A), and polymorphic probes, used to assess genotypes (Figure 1B).¹¹ For copy number analysis, patient DNA labeled with a fluorochrome provides signal intensity that, when compared to a set of reference DNAs, indicates whether there is a

gain or loss of genetic material. For genotype analysis, a single nucleotide polymorphism (SNP) has a single base pair substitution (A, T, C, or G) of one nucleotide to another, then, the alleles corresponding to the nucleotide base changes are arbitrarily given the designation allele A and allele B (polymorphic probes) and reveal which of the genotypes (for example, AA, BB or AB) is present.¹²

In order to better define the genetic abnormalities, the objective of this study was to identify the similarities and differences between SNPa analysis and karyotyping in 25 AML/MDS cases at diagnosis.

Methods

Case selection

Twenty-five patients with AML ($n=22$, non-acute promyelocytic leukemia) and MDS ($n=3$) diagnosed according to WHO criteria at the Escola Paulista de Medicina, Universidade Federal de São Paulo (UNIFESP) from March 2012 to January 2014 were studied.⁴ Bone marrow (BM) aspirate was collected at diagnosis from all patients.

At the same time, buccal cells (BC) were also collected (DNA-SAL™Oasis Diagnostics®) to distinguish clonal abnormalities from constitutional findings.^{13,14}

The study was approved by the Ethics Committees of UNIFESP and Grupo Fleury and was conducted in accordance with the Helsinki Declaration as revised in 2008. All patients gave informed consent prior to entering the study.

Karyotyping

The karyotyping was performed for all BM samples at the time of initial diagnosis and conventional cytogenetic analyses of 24–48 h cultures were performed on bone marrow aspirates using standard techniques. The karyotype results were described according to the International System for Human Cytogenetic Nomenclature 2009.^{15–17} Whenever possible, at least 20 metaphases were analyzed.¹⁷

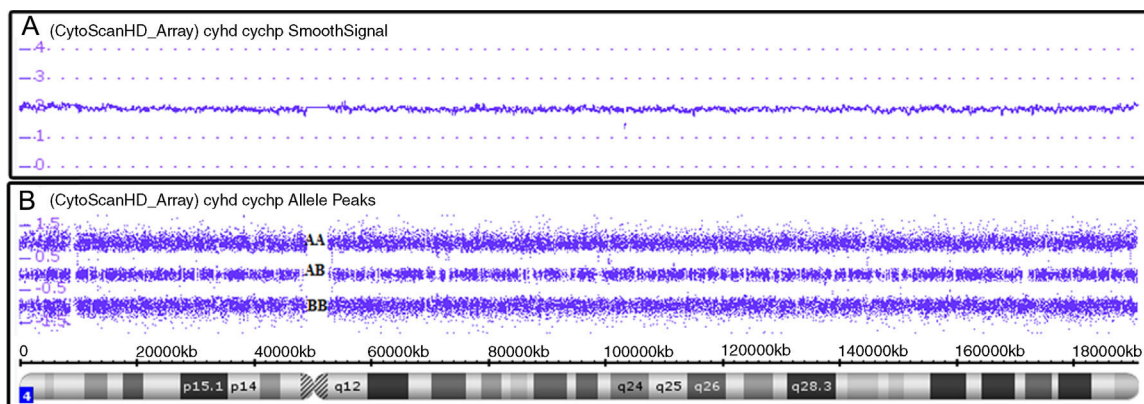


Figure 1 – Affymetrix® Chromosome Analysis Suite image showing normal chromosome 4. (A) Smooth signal representing a normal copy number (purple line CN: 2.00). The information on copy number variations is from non-polymorphic probes. (B) Allele peaks representing a normal genotype with AA, AB and BB alleles (three purple lines). Information on the genotype is from polymorphic probes.

Single nucleotide polymorphism array analysis

SNPa analysis was performed for all samples (BM and BC).¹³ Mononuclear cells from BM were separated (Ficoll method) and DNA was extracted from these cells as well as from BC (Nucleic Acid and Protein Purification kit from Marcherey-Nagel®).^{14,18} DNA (250 ng) was digested, amplified, purified, fragmented, labeled and hybridized using Affymetrix CytoScan® HD Array GeneChip; CEL files were created using the GeneChip® System 3000 7G according to the manufacturer's instructions (Affymetrix). The CEL files were analyzed using Chromosome Analysis Suite v2.0 (ChAS) software.^{19,20}

Equipment, instrumentation, and methodologies employed during the use of microarray platforms were calibrated, monitored, and regularly maintained as appropriate. Positive (Genomic DNA Control supplied by Affymetrix) and negative controls (Low EDTA TE Buffer) were used to validate chip, reagents, and instruments and every CytoScan® HD array has internal quality control metrics used to determine pass/fail for individual samples.²¹

Regions of CNV larger than 1 Mb and CN-LOH larger than 10 Mb identified by the ChAS Software or detected by visual inspection, regardless of gene content, are denoted as true aberrations, with the exception of those known to be normal genomic variants (present in the Genomic Variants Database [<http://projects.tcag.ca/variation>]) and/or found in constitutional (BC) SNPa analysis.^{9,18,22,23}

SNPa results of BC indicate the individual constitutional genome. Thus, changes observed in BM and not found in BC are indicative of acquired abnormalities and were considered for evaluation.¹³

Results

The mean age of the patients was 54 years and the median age was 55 years (range: 28–93). Twelve patients (48%) were male and 13 (52%) were female.

Karyotyping and SNPa results are listed in [Table 1](#).

Karyotype

Ten patients had abnormal karyotypes (40.0%), 11 normal (44.0%) and four had no mitosis (16.0%) ([Figure 2](#)).

Karyotyping detected a total of 17 alterations (8 deletions/losses, 7 trisomies/gains and 2 translocations) in the 25 cases ([Figure 3](#)).

Single nucleotide polymorphism array

By BM SNPa analysis, 17 patients were abnormal (68.0%) and eight normal (32.0%) ([Figure 2](#)) and by BC SNPa analysis, six patients were abnormal (24.0%), 17 were normal (68.0%) and two had no results (8.0%). The internal quality control of Genechip failed for two samples.

SNPa analysis detected a total of 42 alterations (17 losses, 16 gains and 9 CN-LOH) in the 25 cases ([Figure 3](#)).

The SNPa method detected all abnormalities found by karyotype except for:

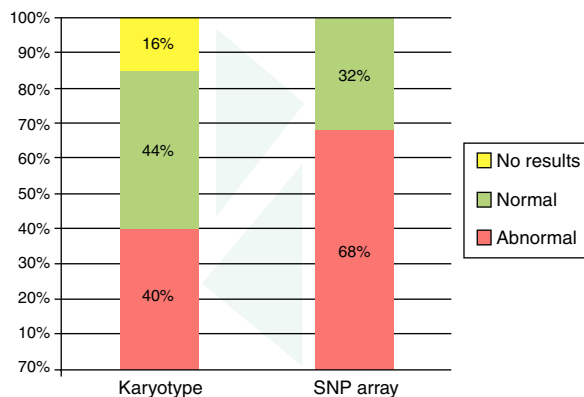


Figure 2 – Results of karyotyping (abnormal: 10 patients, normal: 11 patients and no results: 4 patients) and bone marrow single nucleotide polymorphism array analysis (SNPa; abnormal: 17 patients and normal: 8 patients).

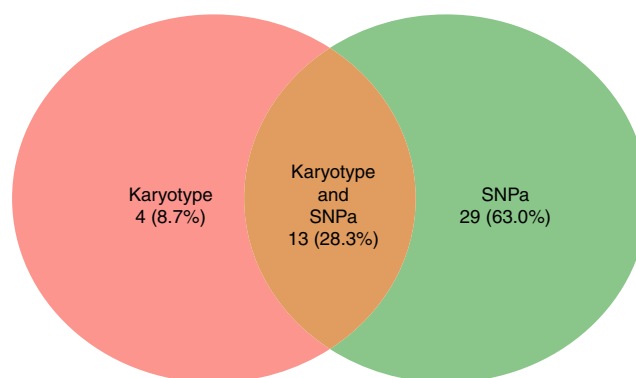


Figure 3 – Frequency of abnormalities detected by karyotyping only: 4 (8.7%), by bone marrow single nucleotide polymorphism array (SNPa) analysis only: 29 (63.0%) and by both methods simultaneously: 13 (28.3%).

- polyploidy in Case 6;
- t(8;21) in Cases 19 and 20;
- ?del(3)(q26) in Case 22.

On the other hand, SNPa detected many abnormalities not found by karyotyping:

- CN-LOH in 3p21.31p21.2, -7, +8, loss in 12p13.33p12.3, 12p12.1p11.22, 12q22q23.3, -16 and -Y in Case 1;
- mosaic gain in 1q31.1q44 and 6q22.33q27, +8 mosaic loss in 2q24.2q37.3, 5q15.q31.1, 5q31.3q32, 5q33.1q35.1 and CN-LOH in 16q21q24.3 in Case 6;
- CN-LOH in 9p24.3p13.3 in Case 13 ([Figure 4A](#));
- +8 in Case 14;
- mosaic gain in 19p13.3p13.11 and 19q13.31q13.33 in Case 17;
- gain in 6q27 in Case 22;
- CN-LOH in 4q31.21q34.3 and 12q23.1q24.11 in Case 23;
- CN-LOH in 6q13q14.3 and 21q21.1q22.3 in Case 24.

The following abnormalities were found by SNPa analysis in cases that karyotyping had no mitosis:

Table 1 – Karyotyping and single nucleotide polymorphism array results.

Case	Diagnosis	Karyotyping	SNPa (BM)	SNPa (BC)
1	AML	46,XY[20]	Normal	Normal
2	AML	46,XY,del(5)(q15q33),del(17)(p11.2)[16]/46,XY[4]	3p21.31p21.2 CN-LOH; 5q21.1q35.3 loss (CN: 1.00); -7; +8; 12p13.33p12.3 loss (CN: 1.00); 12p12.1p11.22 loss (CN: 1.00); 12q22q23.3 loss (CN: 1.00); -16; 17p13.3p11.2 mosaic loss (CN: 1.50); -Y	DGV ^a
3	AML	46,XX[20]	Normal	Normal
4	AML	46,XX[20]	Normal	Normal
5	AML	47,XY,+4[10]	+4	Normal
6	AML	92,XXYY[20]	1q31.1q44 mosaic gain (CN: 2.41); 2q24.2q37.3 mosaic loss (CN: 1.48); 5q15q31.1 mosaic loss (CN: 1.72); 5q31.3q32 mosaic loss (CN: 1.76); 5q33.1q35.1 mosaic loss (CN: 1.73); 6q22.33q27 mosaic gain (CN: 2.31); +8; 16q21q24.3 CN-LOH	Normal
7	AML	47,XX,+8[17]/46,XX[3]	8 mosaic gain (CN: 2.65)	DGV ^a
8	AML	46,XY,del(9)(q21)[20]	9q21.11q22.33 loss (CN: 1.00)	9q21.11q22.33 loss (CN: 1.00)
9	AML	No mitoses	+8; 17q21.2q25.3 CN-LOH	Normal
10	AML	No mitoses	9 mosaic gain (CN: 2.69)	9 mosaic gain (CN: 2.48)
11	AML	No mitoses	Normal	Normal
12	AML	46,XY[15]	Normal	Normal
13	AML	48,XX,+8,+21[11]/46,XX[1]	8 mosaic gain (CN: 2.57); 21 mosaic gain (CN: 2.54); 9p24.3p13.3 CN-LOH	Normal
14	AML	46,XX[10]	+8	8 mosaic gain (CN: 2.93)
15	AML	No mitoses	1q21.1q44 gain (CN: 3.00); 5q15.33q11.2 mosaic gain (CN: 2.39)	Normal
16	AML	46,XX[14]	Normal	Normal
17	AML	46,XX[20]	19p13.3p13.11 mosaic gain (CN: 2.31); 19q13.31q13.33 mosaic gain (CN: 2.22)	Normal
18	AML	46,XY[20]	Normal	Normal
19	AML	46,XY,t(8;21)(q22;q22)[20]/47,XY,+8,t(8;21)(q22;q22)[3]/46,X,-Y,+8,t(8;21)(q22;q22)[2]	+8; -Y	Normal
20	AML	45,X,t(8;21)(q22;q22)[5]/46,XX[5]	X mosaic loss (CN: 1.25)	Normal
21	AML	46,XX[15]	Normal	Normal
22	AML	48,XY,?del(3)(q26),del(11)(q23),+i(21)(q10)x2[cp20]	6q27 gain (CN: 3.00); 11q23.3q25 loss (CN: 1.00); 21q11.2q22.3 gain (CN: 4.75)	Normal
23	MDS	46,XX[20]	4q31.21q34.3 CN-LOH; 12q23.1q24.11 CN-LOH	4q31.21q34.3 CN-LOH; 12q23.1q24.11 CN-LOH
24	MDS	46,XY[15]	6q13q14.3 CN-LOH; 21q21.1q22.3 CN-LOH	6q13q14.3 CN-LOH
25	MDS	46,XXdel(5)(q13q33)[19]/46,XX[1]	5q15q33.2 loss (CN:1.00)	5q15q33.2 loss (CN:1.00)

SNPa: single nucleotide polymorphism array; BM: bone marrow; BC: buccal cells; AML: acute myeloid leukemia; MDS: myelodysplastic syndrome; CN: copy number; LOH: loss of heterozygosity; DGV: genomic variants database.

Loss: CN < 2.00; gain: CN > 2.00; normal: CN = 2.00. Y and X for male: normal: CN = 1.00.

^a BC SNPa (constitutional) was done but failed in the quality control metrics, in these cases the DGV data base was used to validate the findings in BM SNPa test.

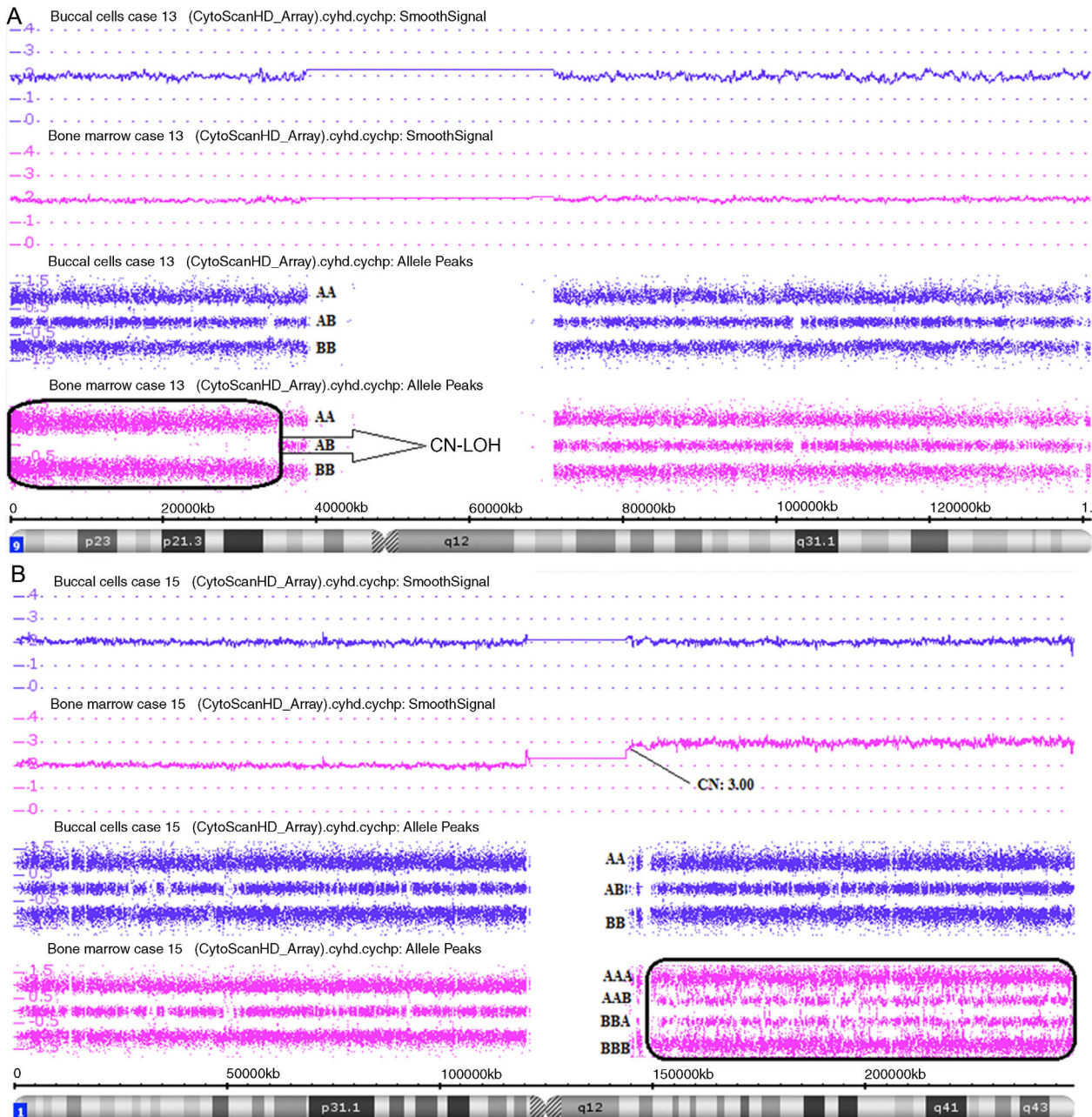


Figure 4 – Affymetrix® Chromosome Analysis Suite Image. (A) Paired analysis (Buccal cell and bone marrow single nucleotide polymorphism array analysis) of chromosome 9 showing copy-neutral loss of heterozygosity (CN-LOH) only in bone marrow. (B) Paired analysis (Buccal cell and bone marrow single nucleotide polymorphism array analysis) of chromosome 1 showing 1q21.1q44 gain only in bone marrow.

- +8 and CN-LOH in 17q21.2q25.3 in Case 9;
- mosaic gain in all chromosome 9 in Case 10;
- gain in 1q21.1q44, mosaic gain in 5q15.33q11.2 in Case 15 (Figure 4B).

Discussion

Cytogenetic abnormalities identified by karyotyping remain the most important prognostic factor in AML.^{2,24}

Molecular karyotyping of these cases by SNPa analysis demonstrated additional information regarding microscopic, submicroscopic, and CN-LOH alterations; the combination of SNPa analysis with karyotyping improves the detection rate for abnormalities.²³

Clearly, the SNPa method does not replace karyotyping as it cannot detect balanced translocations that are relevant to the diagnosis and management of a variety of hematopoietic malignancies.¹¹

In general, cytogenetic abnormalities observed in MDS, unlike AML, are losses or gains of genetic material. Despite

the small number of MDS patients included in this study (three cases), all had abnormalities detected by SNP analysis, while only one had an abnormal karyotype.

CN-LOH regions were found in Cases 6, 9, 13, 23 and 24 (20% of cases). This abnormality represents an important mechanism by which point mutations and other micro lesions can be established in an homozygous state detectable by SNP analysis.^{13,25} CN-LOH is thought to be positively selected during clonal evolution because it can result in homozygosity for a mutation in one allele of a tumor suppressor gene or oncogene, together with the loss of the wild-type allele.^{13,26}

Constitutional SNP analysis (BC) was done in all cases in order to avoid false discoveries by comparing with the findings for BM (Figure 4).^{13,18} In Case 23 two regions of constitutional CN-LOH were found in 4q31.21q34.3 and 12q23.1q24.11. In Case 24 one region of constitutional CN-LOH was found in 6q13q14.3. Therefore, the interpretation of these cases is that the abnormalities were not acquired, but constitutional.

In four cases, the same alteration was found both by BM and BC SNP analysis (constitutional). The alterations found by BC SNP analysis in Cases 10 (mosaic gain in chromosome 9) and 14 (mosaic gain in chromosome 8) were present at a lower level than by the BM SNP analysis, thus suggesting that the BC might have been contaminated with blood at the time of collection, though these changes might have been diluted in BC.¹³

In Cases 8 (loss in 9q21.11q22.33) and 25 (loss in 5q15q33.2), the alterations found in the BC SNP analysis were seen by karyotyping and interpreted as an acquired alteration because in Case 8 this deletion would result in constitutional dimorphism or related syndrome, while in Case 25 the clone was detected by karyotyping along with normal cells. Supporting this interpretation, beyond the scope of this work, we performed SNP analysis in Case 8 after induction therapy and the proportion of the change (CNV) was lower, thereby suggesting that this abnormality was affected by chemotherapy. With this evidence, we consider these two abnormalities as acquired.

The quality control metrics in the constitutional (BC) SNP analysis failed in Cases 2 and 7 and in these cases we used the Genomic Variants Database to validate the findings of the BM SNP analysis.²⁷ Possibly, the failure was due to poor quality DNA from BC.

In Cases 2 and 14, after SNP analysis, some chromosomal abnormalities (-7, +8, -16 and -Y) were found in sporadic metaphases during the karyotyping reanalysis, but had not initially been described because they did not meet the criterion to be considered as a cytogenetic clone,¹⁵ that is, more than three cells with the same numerical change or two cells with the same rearrangement.²⁸

The application of the SNP technique has been largely limited to exploratory research on cancer genetics. However, given its excellent performance in detecting genetic abnormalities in cancer, its application in clinical onco-hematology could be a logical step to attempt to establish better management of cancer patients.^{11,29} The redundancy of tests is beneficial for comprehensive and indistinct characterization of the disease.

SNP analysis may add value to karyotyping-non-informative results and occasionally reveal cryptic abnormalities not recognized by karyotyping.²³ However, SNP analysis should be viewed as a complimentary tool. Some abnormalities detected by SNP analysis and not always recognized by karyotyping are not yet included in classification and prognosis criteria.³⁰

Notwithstanding the progress in this area, SNP tests have not been studied to their full potential and many questions remain to be answered; for instance:

- In which situation could SNP be performed in peripheral blood samples and would the test offer the same information as for marrow samples?
- Do abnormalities found only by SNP analysis present the same prognostic value as karyotyping?
- Should abnormal karyotypes also be investigated by SNP in order to amplify the detection of aberrations?
- Should treatment options targeting specific chromosomal abnormalities be investigated by SNP analysis before introducing the drug?
- Which SNP results would drive a treatment change?
- What additional costs are acceptable in order to obtain better treatment definition?

Conclusion

In summary, SNP analysis was standardized in AML/MDS and results were compared with the abnormalities detected by karyotyping. SNP analysis increased the detection rate of abnormalities compared to karyotyping and also identified a new set of abnormalities that should be further investigated in future studies.

Conflict of interest

The authors declare no conflicts of interest.

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REFERENCES

1. Meyer SC, Levine RL. Translational implications of somatic genomics in acute myeloid leukaemia. *Lancet Oncol.* 2014;15(9):e382-94.
2. Costa AR, Vasudevan A, Krepischi A, Rosenberg C, Chauffaille ML. Single-nucleotide polymorphism-array improves detection rate of genomic alterations in core-binding factor leukemia. *Med Oncol.* 2013;30(2):579.
3. Adès L, Itzykson R, Fenaux P. Myelodysplastic syndromes. *Lancet.* 2014;383(9936):2239-52.
4. Döhner H, Estey EH, Amadori S, Appelbaum FR, Büchner T, Burnett AK, et al. Diagnosis and management of acute

- myeloid leukemia in adults: recommendations from an international expert panel, on behalf of the European LeukemiaNet. *Blood*. 2010;115(3):453-74.
5. Tiu RV, Visconte V, Traina F, Schwandt A, Maciejewski JP. Updates in cytogenetics and molecular markers in MDS. *Curr Hematol Malig Rep*. 2011;6(2):126-35.
 6. Welch JS, Ley TJ, Link DC, Miller CA, Larson DE, Koboldt DC, et al. The origin and evolution of mutations in acute myeloid leukemia. *Cell*. 2012;150(2):264-78.
 7. Raghavan M, Lillington D, Skoulakis S, Debernardi S, Chaplin T, Foot NJ, et al. Genome-wide single nucleotide polymorphism analysis reveals frequent partial uniparental disomy due to somatic recombination in acute myeloid leukemias. *Cancer Res*. 2005;65(2):375-8. Available from: <http://cancerres.aacrjournals.org/content/65/2/375.short>
 8. Frohling S. Comparison of cytogenetic and molecular cytogenetic detection of chromosome abnormalities in 240 consecutive adult patients with acute myeloid leukemia. *J Clin Oncol*. 2002;20(10):2480-5.
 9. Simons A, Sikkema-Raddatz B, de Leeuw N, Konrad NC, Hastings RJ, Schoumans J. Genome-wide arrays in routine diagnostics of hematological malignancies. *Hum Mutat*. 2012;33(6):941-8.
 10. Speicher M, Carter N. The new cytogenetics: blurring the boundaries with molecular biology. *Nat Rev Genet*. 2005;6(10):782-92.
 11. Sato-Otsubo A, Sanada M, Ogawa S. Single-nucleotide polymorphism array karyotyping in clinical practice: where, when, and how? *Semin Oncol*. 2012;39(1):13-25.
 12. Schwartz S. Clinical utility of single nucleotide polymorphism arrays. *Clin Lab Med*. 2011;31(4):581-94, viii.
 13. Heinrichs S, Li C, Look AT. SNP array analysis in hematologic malignancies: avoiding false discoveries. *Blood*. 2010;115(21):4157-61.
 14. de Leeuw N, Hehir-Kwa JY, Simons A, Geurts van Kessel A, Smeets DF, Faas BH, et al. SNP array analysis in constitutional and cancer genome diagnostics—copy number variants, genotyping and quality control. *Cytogenet Genome Res*. 2011;212-21.
 15. Simons A, Shaffer LG, Hastings RJ. Cytogenetic Nomenclature: Changes in the ISCN 2013 Compared to the 2009 Edition. *Cytogenet Genome Res*. 2013 [Epub ahead of print].
 16. Yi JH, Huh J, Kim HJ, Kim SH, Kim SH, Kim KH, et al. Genome-wide single-nucleotide polymorphism array-based karyotyping in myelodysplastic syndrome and chronic myelomonocytic leukemia and its impact on treatment outcomes following decitabine treatment. *Ann Hematol*. 2013;45:9-69.
 17. Pinheiro R, Chauffaille M. Comparison of I-FISH and G-banding for the detection of chromosomal abnormalities during the evolution of myelodysplastic syndrome. *Braz J Med Biol Res*. 2009;42(11):425.
 18. Parkin B, Ouillette P, Li Y, Keller J, Lam C, Roulston D, et al. Clonal evolution and devolution after chemotherapy in adult acute myelogenous leukemia. *Blood*. 2013;121(2):369-77.
 19. Yamamoto G, Nannya Y, Kato M, Sanada M, Levine RL, Kawamata N, et al. Highly sensitive method for genomewide detection of allelic composition in nonpaired, primary tumor specimens by use of affymetrix single-nucleotide-polymorphism genotyping microarrays. *Am J Hum Genet*. 2007;81(1):114-26.
 20. Huggins R, Li LH, Lin YC, Yu AL, Yang HC. Nonparametric estimation of LOH using Affymetrix SNP genotyping arrays for unpaired samples. *J Hum Genet*. 2008;53(11-12):983-90.
 21. Cooley LD, Lebo M, Li MM, Slovak ML, Wolff DJ, Working Group of the American College of Medical Genetics and Genomics (ACMG) Laboratory Quality Assurance Committee. American College of Medical Genetics and Genomics technical standards and guidelines: microarray analysis for chromosome abnormalities in neoplastic disorders. *Genet Med*. 2013;15(6):484-94.
 22. Gibson SE, Luo J, Sathanoori M, Liao J, Surti U, Swerdlow SH. Whole-genome single nucleotide polymorphism array analysis is complementary to classical cytogenetic analysis in the evaluation of lymphoid proliferations. *Am J Clin Pathol*. 2014;141(2):247-55.
 23. Tiu RV, Gondek LP, O'Keefe CL, Huh J, Sekeres MA, Elson P, et al. New lesions detected by single nucleotide polymorphism array-based chromosomal analysis have important clinical impact in acute myeloid leukemia. *J Clin Oncol*. 2009;27(31):5219-26.
 24. Costa AR, Belangero SI, Melaragno MI, Chauffaille ML. Additional chromosomal abnormalities detected by array comparative genomic hybridization in AML. *Med Oncol*. 2012;29(3):2083-7.
 25. Xu X, Johnson EB, Levertson L, Arthur A, Watson Q, Chang FL, et al. The advantage of using SNP array in clinical testing for hematological malignancies: a comparative study of three genetic testing methods. *Cancer Genet*. 2013;206(9-10):317-26.
 26. Radtke I, Mullighan CG, Ishii M, Sua X, Chenga J, Mab J, et al. Genomic analysis reveals few genetic alterations in pediatric acute myeloid leukemia. *Proc Natl Acad Sci U S A*. 2009;106(31). Available from: <http://www.pnas.org/content/106/31/12944.short>
 27. Milosevic JD, Puda A, Malcovati L, Berg T, Hofbauer M, Stukalov A, et al. Clinical significance of genetic aberrations in secondary acute myeloid leukemia. *Am J Hematol*. 2012;87(11):1010-6.
 28. Chauffaille ML. Alterações cromossômicas em síndrome mielodisplásica. *Rev Bras Hematol Hemoter*. 2006;28(3):182-7.
 29. Yi JH, Huh J, Kim HJ, Kim SH, Kim HJ, Kim YK, et al. Adverse prognostic impact of abnormal lesions detected by genome-wide single nucleotide polymorphism array-based karyotyping analysis in acute myeloid leukemia with normal karyotype. *J Clin Oncol*. 2011;29(35):4702-8.
 30. Chauffaille ML. Adding FISH to karyotype in myelodysplastic syndrome investigation diagnosis: are all questions answered? *Leuk Res*. 2010;34(4):413-5.