Prevalence and clinical implications of germline predisposition gene mutations in patients with acute myeloid leukemia

Borahm Kim1,2, Woobin Yun3, Seung-Tae Lee2, Jong Rok Choi2, Keon Hee Yoo4, Hong Hoe Koo4, Chul Won Jung5 & Sun Hee Kim6

Acute myeloid leukemia (AML) is one of the most common types of leukemia. With the recent advances in sequencing technology and the growing body of knowledge on the genetics of AML, there is increasing concern about cancer predisposing germline mutations as well as somatic mutations. As familial cases sharing germline mutations are constantly reported, germline predisposition gene mutations in patients with AML are gaining attention. We performed genomic sequencing of Korean patients diagnosed with AML to identify the prevalence and characteristics of germline predisposition mutations. Among 180 patients, germline predisposition mutations were identified in 13 patients (13/180, 7.2%, eight adults and five children). Germline mutations of BLM, BRCA1, BRCA2, CTC1, DDX41, ERCC4, ERCC6, FANCI, FANCM, PALB2, and SBDS were identified. Most of the mutations are in genes involved in DNA repair and genomic stability maintenance. Patients harboring germline mutations tended to have earlier onset of AML (p = 0.005), however, the presence of germline mutations did not show significant association with other clinical characteristics or treatment outcome. Since each mutation was rare, further study with a larger number of cases would be needed to establish the effect of the mutations.

With the recent advances in sequencing technology and the growing body of knowledge on the genetics of AML, there is increasing concern about cancer predisposing germline mutations as well as somatic mutations. It has been widely recognized that not only somatic mutations in cancer tissue, but also germline gene mutations can affect disease characteristics, progress, and prognosis, and in cases with solid cancers such as breast cancer, the significance of germline mutation had been already recognized and changes in treatment and genetic counseling according to the presence of those mutations had been settled down in clinical practice. Similarly, the category myeloid neoplasm with germline predisposition mutations was included in the WHO classification in 2016. A number of cases of familial leukemia with these mutations was reported, mainly in relatively well-recognized genes such as DDX41, CEBPA, and RUNX1. However, these studies focused on specific ethnic groups, and data regarding other ethnicities are lacking. Furthermore, for patients with AML, hematopoietic stem cell transplantation (HSCT) is frequently performed. Germline predisposition mutations could be a significant issue in the setting of HSCT, where most donors are family members, and a higher probability of shared mutations is expected. As in the case of BRCA1/2 gene mutation, there may be an increased risk of other cancers, and patients and family members having the same mutation should be involved in a surveillance program. Therefore, there is a growing need for basic information about frequency and types of germline predisposition gene mutations.

In this context, we assessed the prevalence of germline predisposition gene mutations and identified the clinical characteristics of mutation carriers among Korean patients with AML using genomic sequencing and

1Department of Laboratory Medicine, Korea University College of Medicine, Seoul, Korea. 2Department of Laboratory Medicine, Yonsei University College of Medicine, 50 Yonsei-ro, Seodaemun-gu, Seoul 03722, Korea. 3Brain Korea 21 PLUS Project for Medical Science, Yonsei University, Seoul, Korea. 4Department of Pediatrics, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul, Korea. 5Department of Internal Medicine, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul, Korea. 6Department of Laboratory Medicine and Genetics, Samsung Medical Center, Sungkyunkwan University School of Medicine, 81 Ilwon-ro, Gangnam-gu, Seoul 06351, Korea. *email: lee.st@yuhs.ac; sunnyhk@skku.edu
we correlated the mutation status with treatment outcome, to provide a broader insight into AML biology and patient care.

**Methods**

**Patients.** Bone marrow aspirates of 180 patients diagnosed with AML in our center between 2013 and 2017 were obtained after receiving informed consent for genetic study from each patient. This study was approved by the Samsung Medical Center institutional review board (2017-04-027) and was conducted in accordance with the tenets of the Declaration of Helsinki.

**Genomic sequencing.** A total of 139 patients was evaluated by whole exome sequencing (WES) and 41 patients were by gene panel sequencing. NGS testing was performed with initial diagnostic samples. Clinical characteristics of patients were in Supplemental Table 1. For WES, libraries were prepared using the Twist Human Core Exome Kit (Twist Bioscience, San Francisco, CA, USA) and sequenced on a NovaSeq system (Illumina, San Diego, CA, USA). For gene panel sequencing, gene panel containing 215 genes (Supplemental Table 2) associated with hematologic malignancy or germline cancer predisposition was used and NextSeq 550 instrument (Illumina) was used. Detailed sequencing methods and bioinformatics analysis were in Supplemental Methods.

**Interpretation of variants.** The following databases were used for variant annotation: Online Mendelian Inheritance in Man (OMIM), Human Gene Mutation Database (HGMD), ClinVar, dbSNP, 1,000 Genome, Genome Aggregation Database (gnomAD), Exome Sequencing Project (ESP), and the Korean Reference Genome Database (KRGDB). The pathogenicity of missense variants was predicted using five in silico prediction algorithms, including Sorting Tolerant from Intolerant (SIFT), Polymorphism Phenotyping v2 (PolyPhen-2), MutationTaster, MutationAssessor, and Functional Analysis through Hidden Markov Models (FATHMM), implemented in dbNSFP. Effects on splicing were predicted using SPIDEX, and dbSNV.

We first established a set of genes known to be associated with AML predisposition according to WHO classification, and variants of those genes were prioritized. Variants were further classified according to American College of Medical Genetics (ACMG) guideline. For PM2 score, global population frequency cut off < 0.00001 for dominant disease and < 0.0001 for recessive disease were applied. For PP3 score, the agreement of at least five prediction tools was applied.

**Germline confirmation test.** Suspected variants in germline predisposition genes were further confirmed with bone marrow specimens were collected when the patients were in complete remission. Sanger sequencing was performed using custom primers and the BigDye Terminator Cycle Sequencing Ready Reaction Kit on an ABI Prism 3730 Genetic Analyzer (Applied Biosystems, Foster City, CA, USA). Because bone marrow was used for germline mutation testing, possibility of confounding due to factors including residual tumor and clonal hematopoiesis cannot be ruled out. Interpretation of Sanger sequencing results was proceeded under recognition of the limitation.

**Statistical analysis.** To compare the outcome according to molecular characteristics, Fisher’s exact test and logistic regression analysis were utilized, with single or multiple variables. Variables included in multiple logistic levels were chosen by adapting stepwise as a variable selection method. Statistical analyses were computed using R. *p* values < 0.05 were considered statistically significant.

**Results/discussion**

For gene panel sequencing, diagnostic exome sequencing, median sequence coverage was 1626x, with an average of genotype quality score 98. For whole exome sequencing, median sequence coverage was 182x, with an average of genotype quality score 54.

**Genes involved in DNA repair or maintenance of genomic stability were frequently mutated.** Among 180 patients, 13 (13/180, 7.2%) showed pathogenic mutations in germline predisposition genes (Table 1). Most of the identified germline predisposition gene mutations were in genes involved in DNA repair or maintenance of genomic stability, which were associated with inherited bone marrow failure syndromes including Fanconi anemia (FA), dyskeratosis congenita (DC), or Shwachman–Diamond syndrome (SDS).

Eight of the 13 germline mutations identified were in six FA genes: *FANCD1 (BRCA2), FANCI, FANCM, FANCN (PALB2), FANQC (ERCC4)*, and *FANCS (BRCA1)*. FA genes are involved in the FA signaling pathway, which is crucial in the DNA damage response. Constant exposure to endogenous and exogenous genotoxic agents can jeopardize genomic stability when the DNA damage response is compromised. FA is a rare genetic disorder often accompanied by numerous other conditions, including early age of onset of symptoms, multiorgan congenital defects, bone marrow failure leading to pancytopenia, and predisposition to hematological and non-hematological malignancies. Monoallelic mutation carriers who had no apparent signs, were known to have increased risk for cancer. The protein products of these six genes are components of the DNA damage response system and participate in various cellular processes. Genomic instability resulting from defective function of those proteins can be associated with increased cancer risk. However, elucidation of the function of each component and the consequences of deficiencies of each product have yet to be determined.

In addition, *BRCA1/2* and *PALB2* are not only associated with FA and predisposition for AML, but are also widely known as important risk factors for breast and ovarian cancer. Although monoallelic mutation carriers
| ID     | Age | Gene     | OMIM disease                                         | Variant\(^a\) | VAF  | Depth | Type       | ClinVar                  | ACMG classification                  | Population frequency\(^b\) | Accompanied somatic mutations\(^c\) |
|--------|-----|----------|------------------------------------------------------|---------------|------|-------|------------|--------------------------|-------------------------------|-------------------------------|-----------------------------------|
| P1561  | 30  | BRCA1    | Familial breast-ovarian cancer 1, 404,370, AD        | c.2433delC    | 0.54 | 127   | Frameshift | Pathogenic              | Pathogenic (PM2, PP3)           | 0.00005437                      | KMT2A-MLLT3                       |
|        |     | NM_007294.3 |                                      | p.Lys812ArgfsTer3 |     |       |            |                          |                               |                                |                                   |
| P3065  | 62  | BRCA2    | Familial breast-ovarian cancer 2, 412,555, AD        | c.2798_2799delCA| 0.39 | 137   | Frameshift | Pathogenic              | Pathogenic (PM1, PM2, PP3)      | 0                              | IDH2, DNMT3A                      |
|        |     | NM_000059.3 | Fanconi anemia, complementation group D1, 605,724, AR | p.Thr933ArgfsTer2 |     |       |            |                          |                               |                                |                                   |
| P1052  | 45  | PALB2    | Fanconi anemia, complementation group N, 610,832     | exon 11 deletion | 0.51 | 1,050 | Frameshift | Not reported            | Likely pathogenic (PM2)          | 0                              | DNMT3A, FLT3 ITD, KMT2A PTD      |
|        |     | NM_024675.3 | Breast cancer, susceptibility to, 114,480, AD        |               |      |       |            |                          |                               |                                |                                   |
|        |     |          | Pancreatic cancer, susceptibility to                  |               |      |       |            |                          |                               |                                |                                   |
| P1257  | 7   | FANCM    | Premature ovarian failure 15, 618,096, AR            | c.1456C>T     | 0.54 | 254   | Nonsense   | Not reported            | Likely pathogenic (PM2)          | 0                              | BRCA2                            |
|        |     | NM_020937.2 | Spermatogenic failure 28, 618,086, AR             | p.Arg486Ter   |      |       |            |                          |                               |                                |                                   |
| P2912  | 58  | FANCI    | Fanconi anemia, complementation group I, 609,053     | c.158-2A>G    | 0.54 | 113   | Splice site | Likely pathogenic     | Likely pathogenic (PM2)          | 0.00005437                      | RUNXI-RUNXI1                      |
|        |     | NM_001113378.1 | Fanconi anemia, complementation group I, 609,053 |               |      |       |            |                          |                               |                                |                                   |
| P1415  | 3   | FANCI    | Fanconi anemia, complementation group I, 609,053     | c.158-2A>G    | 0.6  | 115   | Splice site | Likely pathogenic     | Likely pathogenic (PM2)          | 0.00005437                      | KIT                               |
|        |     | NM_001113378.1 | Fanconi anemia, complementation group I, 609,053 |               |      |       |            |                          |                               |                                |                                   |
| P2605  | 5   | ERCC4    | Fanconi anemia, complementation group Q, 615,272, AR | c.458G>A p.Arg153His | 0.48 | 2,311 | Missense   | Not reported            | Likely pathogenic (PM1, PM2, PM5, PP3) | 0.0001631                      | FLT3, FLT3 ITD, PML-RARA          |
|        |     | NM_005236.2 | Aplastic anemia, susceptibility to, 609,135         |               |      |       |            |                          |                               |                                |                                   |
|        |     |          | Shwachman-Diamond syndrome, 260,400, AR              |               |      |       |            |                          |                               |                                |                                   |
| P0655  | 0   | SBDS     | Aplastic anemia, susceptibility to, 609,135         | c.258 + 2T>C  | 0.53 | 90    | Splice site | Pathogenic              | Pathogenic (PM2, PP3)           | 0.009412                       | GATA1                             |
|        |     | NM_016038.2 | Shwachman-Diamond syndrome, 260,400, AR            |               |      |       |            |                          |                               |                                |                                   |
do not develop FA, monoallelic mutations in BRCA1/2, or PALB2 are associated with increased risk of specific cancers, e.g., breast cancer. Recent National comprehensive cancer network (NCCN) guidelines suggest that carriers of those mutations should be included in the surveillance program and should consider preventive actions [https://www.nccn.orgprofessionals/physician_gls/pdf/genetics_screening.pdf]. The same principle should hold for the patient with AML with germline predisposition mutations.

Besides, BLM is associated with Bloom syndrome, a rare chromosomal instability disorder characterized by growth retardation, immunodeficiency, and a wide spectrum of cancers12. BLM mutation is known to be associated with cancers13. ERCC6 plays a critical role in DNA repair, and an association between disruption of its function and increased susceptibility to cancer has been reported14. CTC1 is known causative genes of the telomere biology disorder DC, characterized by accelerated telomere shortening leading to manifestations such as bone marrow failure, cancer, and pulmonary fibrosis15. DC genes function in telomere maintenance; CTC1 functions in a telomere-associated complex to protect the telomere from lethal DNA degradation16. Although these DC genes have autosomal recessive inheritance, an association between monoallelic deleterious germline mutations and myeloid malignancies has been reported17.

Splice site mutations of the SBDS, a causative gene of SDS, were identified in two patients. This mutation also had relatively high frequency among the control population (0.0048 from KRGDB and 0.004 from gnomAD East Asians), probably reflecting high carrier frequency. SBDS is characterized by exocrine pancreatic insufficiency, skeletal abnormalities, and bone marrow failure with an increased risk of myeloid malignancy18. Approximately

Table 1. Identified germline predisposition gene mutations. VAF variant allele fraction, ITD internal tandem duplication, PTD partial tandem duplication, VOUS a variant of unknown significance. Variants were confirmed as germline origin using bone marrow aspirate in complete remission. Population frequencies from gnomAD (https://gnomad.broadinstitute.org/). Tier 1 or Tier 2 somatic variants according to published guideline (Li et al., J Mol Diagn 2017, PMID 27993330) were presented.
The most frequently reported mutation in Caucasians, monoallelic mutation and increased risk of malignancy cannot be excluded. Detected in two of our patients. Although monoallelic mutations are not known to cause SDS, association between patients with germline mutations in other genes such as CEBPA positive group.

which are well-known poor prognostic factors, were not identified in the germline predisposition mutation.

tions was not statistically clear. In addition, somatic mutations of this younger age group. with better outcome > The effect of germline predisposition gene mutations needs to be further investigated in acute lymphoblastic leukemia, it cannot be ruled out harboring germline predisposition mutation was associated the negative finding, the number of identified mutations was possibly insufficient to determine the effects of

90% of SDS is caused by two common mutations, c.183_184delinsCT and c.258 + 2T > G19, only the latter was detected in two of our patients. Although monoallelic mutations are not known to cause SDS, association between monoallelic mutation and increased risk of malignancy cannot be excluded.

The DDX41 mutations were known to have different spectrum depending on ethnic groups. p.A500Cfs*9 mutation has been solely reported in Asian patients. The most frequently reported mutation in Caucasians, p.D140Gs*2, was not found in our patients. The germline DDX41 mutation is one of most frequently detected germline predisposition mutations in myeloid malignancy, with around 70 families described to date. In these families, myeloid malignancies were associated with normal karyotypes, and about 50% were found to have a somatic second hit mutation in DDX41, suggesting that DDX41 acts as a tumor suppressor. DDX41 is associated with the dominant inheritance and donor cell derived leukemia is already reported, screening of germline predisposition mutation should be considered during donor selection.

Germline predisposition mutations were associated with earlier age of onset. Characteristics of patients with germline predisposition mutations are given in Table 2. There was a significant difference in age of onset between the two groups. Patients with germline predisposition mutations tended to be younger, showing an earlier age of onset (p = 0.005). Notably, while only 21 patients under age 20 were included (21/180, 11.7%), five among 13 with germline mutation (5/13, 38.5%) were children. Five of 21 children and eight of 139 adults had germline predisposition mutations (p = 0.005). Although certain mutations like those in DDX41 are reported not to be associated with early onset malignancy, age of AML diagnosis was significantly lower in patients with germline mutations in other genes such as CEBPA. It is understandable that inherited disorders tend to be expressed at earlier age in childhood. < Considering better prognosis of children in patients with acute lymphoblastic leukemia, it cannot be ruled out harboring germline predisposition mutation was associated with better outcome > The effect of germline predisposition gene mutations needs to be further investigated in this younger age group.

Notably, although the association between germline predisposition gene mutations and certain somatic mutations was not statistically clear. In addition, somatic mutations of RUNXI and ASXL1 and complex karyotype, which are well-known poor prognostic factors, were not identified in the germline predisposition mutation positive group.

Presence of germline predisposition gene mutations did not affect the clinical outcome. The presence or absence of germline predisposition mutations, however, did not affect clinical outcome. Factors confirmed as significant in this study were well-recognized good or poor prognostic factors of AML.

In multivariate Cox proportional hazards regression analysis for overall survival (OS), complex karyotype, older age, absence of gene fusion, poor outcome of induction chemotherapy, and FLT3 ITD mutation were factors for unfavorable outcome (Supplemental Table 3). In the same analysis for relapse-free survival (RFS), poor outcome factors were RUNXI somatic mutation and FLT3 ITD (Supplemental Table 4). On the other hand, complete remission after induction chemotherapy, the presence of gene fusions, and CD34-negative immunophenotype mutation were identified as favorable factors. Achievement of complete remission after induction chemotherapy and carrying well-known poor prognostic features like FLT3 ITD mutation were important factors for both OS and RFS.

The presence or absence of germline predisposition mutations did not affect clinical outcome. As reason for the negative finding, the number of identified mutations was possibly insufficient to determine the effects of

| Genetics                  | Median age | OR         | p*          |
|---------------------------|------------|------------|-------------|
| FLT3 ITD                  | 40 (range, 0–62) | 0.941     | 0.96 (0.93–0.99) |
| Gene fusion               | 27 (16.2) | 0.94 (0.14–3.77) |
| No. of somatic mutations  | 1–5       | 0.585    | 1.09 (0.79–1.44) |

Table 2. Characteristics of patients with germline predisposition mutation. OR odds ratio, ITD internal tandem duplication, Number, IP immunophenotype, CR complete remission, HSCT hematopoietic stem cell transplantation. Fisher’s exact test was used. **P < 0.01
those mutations. More patients and germline mutation carriers would be needed to establish whether germline predisposition mutations are beneficial or harmful. Clinical and therapeutic heterogeneity of patients might also play the role. Comparison of patients between groups with otherwise identical condition would be desirable.

Although younger age could be linked to higher durable intensity of and better response to chemotherapy, the association of germline predisposition mutation and treatment outcome was not definite. Because statistical significance could not be achieved partially due to insufficient sample size, further study with more patients is needed.

In conclusion, we identified 13 patients with germline predisposition mutation among 180 patients with AML. Most of the mutated genes are involved in the DNA repair system, contributing genomic stability. Although the effect of these mutations on clinical outcome, including OS and RFS, was not significant, we confirmed that this group of patients tends to develop AML at a younger age. Since each mutation was rare, further study with a larger number of cases would be needed to establish the effect of the mutations.

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B.K. performed the research, analyzed the data, and wrote the paper. W.Y. performed the experiment and analyzed the data. S.L., J.C. and S.K. designed the study and review the paper. K.Y., H.K., and C.J. obtained and analyzed the patient data.

Competing interests
The authors declare no competing interests.

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Correspondence and requests for materials should be addressed to S.-T.L. or S.H.K.

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