Precursor B cell lymphoid blast crisis of chronic myeloid leukemia with novel chromosomal abnormalities: A case report

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Abstract. Chronic myeloid leukemia (CML) is a clonal hematopoietic stem cell disorder. It is characterized by the presence of the Philadelphia (Ph) chromosome, t(9;22)(q34.1;q11.2), which carries the \textit{BCR-ABL1} fusion gene. Tyrosine kinase inhibitors (TKIs) have markedly changed the treatment approach of CML and have become the first-line agents for almost all CML patients. However, certain patients experience resistance to these medications, which occurs through several mechanisms, including the accumulation of TKI-resistant chromosomal abnormalities. The present study reports a case of a 27-year-old Saudi male with CML receiving TKI treatment, who presented with precursor B-cell lymphoblastic crisis demonstrating the presence of the novel combined chromosomal abnormalities; non-Ph der(22), i(9) and der(20), carrying the \textit{BCR-ABL1} fusion gene. This case report adds to the literature on novel TKI-resistance-conferring chromosomal abnormalities and links them to precursor B-cell lymphoblastic crisis.

Introduction

CML is a myeloproliferative disorder that is characterized by the presence of the reciprocal translocation t(9;22), which forms the Philadelphia (Ph) chromosome. During this translocation, the breakpoint cluster region (BCR) gene at position 22q11.2 is juxtaposed to the c-Abelson (ABL1) gene at 9q34.1, forming the \textit{BCR-ABL1} fusion gene, which encodes a constitutively active tyrosine kinase (TK) protein (1,2). The constitutively active protein is associated with increased levels of erythrocytes, monocytes, megakaryocytes, myelocytes and platelets in the peripheral blood and marked myeloid hyperplasia in the bone marrow (3).

CML presents in one of three phases: Chronic phase, accelerated phase or blast crisis. The latter is of myeloid, lymphoid or mixed-lineage phenotype (4). TKIs have markedly changed the approach to CML management. TKIs have improved patient outcomes to the extent that they are now currently accepted as the first-line agents for nearly all patients with CML, regardless of the phase of the disease. However, certain patients experience resistance to these medications; this occurs through several mechanisms including the accumulation of additional cytogenetic abnormalities, which can confer a survival advantage to the treated myeloid cells. The most common cytogenetic abnormalities include an additional Ph chromosome, trisomy 8 and isochromosome 17q (5,6). Several other less common cytogenetic abnormalities have been reported; however, to the best of our knowledge, those found in the present case have not been previously reported.

Case report

Presentation. The present case involves a 27-year-old Saudi male patient, whose initial presentation was three years prior. At that time, he presented with pallor and abdominal distension. He was revealed to have significant splenomegaly and marked leukocytosis, with a white blood cell (WBC) count of 105x10⁹/l. Subsequent investigations confirmed the diagnosis of CML. The patient was initially treated with imatinib; however, due to myelosuppression, the treatment was changed to dasatinib. Subsequently, due to a skin reaction, the treatment was changed to nilotinib (100 mg/day), which the patient clinically responded to and tolerated well.
At his current presentation, the patient had fever, bone pain and cytopenia. Investigations confirmed the diagnosis of precursor B cell acute lymphoblastic leukemia with the presence of the novel combined chromosomal abnormalities of non-Ph der(22), i(9), and der(20), carrying the *BCR-ABL1* fusion gene.

**Complete blood count (CBC).** CBC with differential was performed using an Automatic Hematological Analyzer Sysmex XE-5000 (Sysmex America, Inc., Lincolnshire, IL, USA).

**Immunophenotyping.** Immunophenotyping was performed on the patient's bone marrow aspirate as follows; upon collection of the bone marrow aspirate, 0.5 ml of the sample was mixed with 10 ml of red blood cell lysing solution and centrifuged at 540 x g for 5 min. The supernatant was discarded and the cell pellet further washed with PBS. A 100 µl aliquot of cell suspension with an adjusted concentration of 10x10⁷ cells/ml was added to tubes containing commercial pre-titrated volumes of labelled monoclonal antibody cocktails to bind several antigens, surface and cytoplasmic clusters of differentiation (CD) (BD Biosciences, San Jose, CA, USA) and incubated in the dark for 15 min at room temperature. These monoclonal antibodies were used in conjunction with four fluorochromes [fluorescein isothiocyanate (FITC), phycoerythrin (PE), allophycocyanin (APC) and peridinin chlorophyll (PerCP)] in each tube where they were diluted by factor of 20. FITC labelled antibodies bound to CD14 (cat. no. 561712), surface and cytoplasmic IgG (cat. no. 560952), surface and cytoplasmic CD3 (cat. no. 561806), CD34 (cat. no. 560942), cytoplasmic CD66 (cat. no. 551479), IgM (cat. no. 562029), CD33 (cat. no. 561818), CD38 (cat. no. 560982), CD2 (cat. no. 561759), CD64 (cat. no. 560970), terminal deoxynucleotidyl transferase (TdT) (cat. no. 347194), CD45 (cat. no. 560976) and (myeloperoxidase) MPO (cat. no. 340580). PE labelled antibodies bound to surface and cytoplasmic IgG (cat. no. 560951), cytoplasmic CD11 (cat. no. 560999), CD8 (cat. no. 560950), CD10 (cat. no. 561002), CD56 (cat. no. 561903), CD117 (cat. no. 561682), CD13 (cat. no. 560998), CD7 (cat. no. 561934), CD58 (cat. no. 560959) and cytoplasmic CD79a (cat. no. 555935). APC labelled antibodies bound surface and cytoplasmic IgG (cat. no. 562025), CD20 (cat. no. 560900), CD4 (cat. no. 561840), CD19 (cat. no. 561742), CD15 (cat. no. 561716), CD22 (cat. no. 562860), CD11b (cat. no. 340937), CD5 (cat. no. 340583), HLA-DR (cat. no. 560896), cytoplasmic CD22 (cat. no. 562860), CD25 (cat. no. 340939) and CD34 (cat. no. 560940). PerCP labelled antibodies bound CD45 (cat. no. 561086) and cytoplasmic CD3 (cat. no. 347344).

For intracellular staining (i.e., for CD79a, CD3, TdT and MPO), lymphocyte permeabilization preceded the addition of cytoplasmic and nuclear antibodies. This was achieved by adding 0.5 ml FACs Permeabilizing Solution to 100 µl of the lysed sample followed by 10-minute dark incubation at room temperature. The FACs Permeabilizing Solution was prepared by diluting 1 ml of BD Permeabilizing stock solution (cat. no. 340973; BD Biosciences) in 9 ml of distilled water.

The samples were processed with a FACSCanto II cell analyzer and the analysis was performed using FACs Diva Software (version 8.0.1; BD Biosciences). The flow cytometry data was analyzed with a threshold of 25,000 events. For gating, forward scatter, side scatter, CD45, CD3, CD19 in addition to other lineage specific markers were used. The lineages of the blasts were determined in each case depending upon the expression of lineage-specific markers where an expression for a certain marker was considered positive if the percentage of the cells expressing that marker was ≥20% and negative if <20%, except for TdT and MPO where the threshold was 10% as recommended (7).

**Cytogenetic analysis.** Chromosomal analysis using GTG banding was performed as described previously (8). Karyotyping was performed in 20 metaphases from the patient's unstimulated bone marrow sample according to the nomenclature of the International System for Human Cytogenetics (9).

**Fluorescence in situ hybridization (FISH).** FISH was performed using Vysis *BCR-ABL1* Tri-Color Dual-Fusion FISH Probes (Abbott Pharmaceutical Co. Ltd., Lake Bluff, IL, USA) to detect the *BCR-ABL1* translocation, as described previously (10).

**Molecular analysis.** EDTA whole blood samples from the patient were used for quantification of the *BCR-ABL1* P210 transcript by reverse transcription-quantitative polymerase chain reaction (RT-qPCR). This was performed using the GeneXpert® Dx System (Roche Diagnostics GmbH, Mannheim, Germany) as described previously (11).

**Initial presentation.** At initial presentation, the patient's CBC with differential revealed a hemoglobin (Hb) level of 8.6 g/dl, a platelet count of 188x10⁹/l and a WBC count of 105x10⁹/l, comprised of 52% neutrophils, 1% lymphocytes, 3% monocytes, 0% eosinophils, 0% basophils, 1% promyelocytes, 2% myelocytes and 5% blast cells. The patient's peripheral blood smear revealed normocytic hypochromic anemia with anisocytosis and schistocytosis. There was marked leukocytosis with a significant number of immature myeloid precursors, indicating leukoerythroblasticosis. The patient's bone marrow aspirate was hemodiluted, but revealed moderate cellularity with the presence of myeloid, erythroid and megakaryocytic lineages and ~5% blast cells. The karyotype of each of the 10 metaphases obtained from the bone marrow aspirate was 46,XY,t(9;22)(q34.1;q11.2). FISH analysis confirmed the presence of the *BCR-ABL1* fusion in 95% of nuclei in the aspirate. Molecular studies of the patient's peripheral blood determined a *BCR-ABL1* fusion transcript level of 100% International Scale (IS) units. Based on these results, the patient was diagnosed with CML.

**Current status.** At the current presentation, the patient's CBC with differential revealed an Hb level of 10.5 g/dl, a platelet count of 37x10⁹/l and a WBC count of 0.9x10⁹/l, comprised of 48% neutrophils, 25% lymphocytes, 6% variant lymphocytes, 2% monocytes, 0% eosinophils, 0% basophils and 14% blast cells. The peripheral blood smear revealed pancytopenia with the presence of blasts.
Figure 1. (A) Giemsa-banding karyogram of the patient's bone marrow. Isochromosome 9q, derivative chromosomes 20 and 22 (red arrows) are presented, where the extra material on 20 containing BCR-ABL1 fusion is derived from chromosome 22. (B) Ideogram of all chromosomal abnormalities observed in this case with breakpoints marked.

Figure 2. FISH analysis of one metaphase and one interphase nucleus. (A) Showing two ABL1 (red) signals, one BCR signal (green), and two fusion gene BCR-ABL1 signals (yellow). (B) Labelled inverted image showing two ABL1 signals on the two long arms of isochromosome 9 (red), one BCR signal (green) on chromosome 22, and two fusion gene BCR-ABL1 signals (yellow) one on the long arm of der 9 and one on that of der 20, confirming the abnormalities seen on the karyogram. (C) Non-labelled inverted image identifying the chromosomes containing the signals.
The bone marrow aspirate revealed markedly increased blasts with markedly decreased myeloid, erythroid and megakaryocytic cell lineages. Immunophenotyping analysis of the aspirate revealed that the blast cells were positive for CD34 (partial, i.e., only a subset of the population of interest was positive), cytoplasmic CD79a, CD19, CD10, cytoplasmic TdT, CD20, HLA-DR, CD73 (partial), CD58, CD44, CD200, CD24, cytoplasmic CD66 and CD72 antigens. The cells were negative for cytoplasmic myeloperoxidase, cytoplasmic CD3, surface CD3, CD7, surface IgM, CD45 (negative to low), CD117, cytoplasmic CD22, CD33, CD13, CD38, CD123 and CD86 antigens, consistent with the precursor B-cell lymphoblastic nature of these blast cells.

A bone marrow biopsy revealed 99% cellularity with 95-99% blasts. The number of morphologically normal myeloid, erythroid and megakaryocytic cells was markedly decreased to absent. Karyotyping performed on 20 metaphases from the bone marrow revealed the following: 46,XY,i(9)(q10),der(22)t(9;22)q34.1;q11.2,46,XY[2]/46,XY[t(20;22)(q11.2;q11.2)][18]. This result, combined with that of the FISH analysis, confirmed the presence of a clone with the concurrent cytogenetic abnormalities of i(9)(q10), non-Philadelphia der(22) and der(20) carrying the BCR-ABL1 fusion gene (Figs. 1B and 2A-C).

Discussion

The blast crisis demonstrated in this case was of the precursor B cell lymphoblastic type. It is well established that ~30% of blast crises in CML are of the lymphoid rather than the
myeloid phenotype. The aberrant cytogenetic abnormalities observed in this case, in addition to the BCR-ABL1 fusion during Ph chromosome formation, were i(9q)(q10) formation and the reciprocal translocation between the Ph chromosome and 20q11.2. To the best of our knowledge, this is the first case reported to combine these cytogenetic aberrations. This case report also links these cytogenetic aberrations to the precursor B cell lymphoblast phenotype. Ascertaining the blast phenotype has its own therapeutic implications, since the treatment protocol of lymphoid blast crisis is different to that of the myeloid type. In other words, linking these abnormalities to lymphoid crisis, in this case, had therapeutic implications since this directed the treatment toward vincristine- and prednisone-based protocols (i.e., lymphoid blast crisis protocol).

From the patient's history, initial karyotyping and FISH results, it may be concluded that these cytogenetic abnormalities did not occur simultaneously. The first abnormality was the classical t(9;22)(q34.1;q11.2), resulting in Ph chromosome formation (Fig. 3A). The other two abnormalities were the i(9q)(q10) formation and the reciprocal translocation between Ph and 22 chromosomes, transferring the BCR-ABL1 fusion gene to the latter (Fig. 3B and C). These two events must have followed the Ph chromosome formation during the clonal evolution that conferred TKI resistance. It is not possible to tell whether these two latter events occurred simultaneously, or their order of occurrence if they did occur sequentially. For demonstration purposes only, i(9)(q10) formation is indicated in Fig. 3C to be the final event.

It is likely that the B cell lymphoid nature of the blast crisis observed in this patient is attributed to the loss of 9p during i(9q) formation. This possibility stems from the fact that 9p loss is a known recurrent cytogenetic abnormality observed in chronic lymphocytic leukemia and mantle cell lymphoma (12). This arm contains genes that are important for B cell differentiation and cell cycle regulation, including PAX5 and CDKN2A. The paired box gene PAX5 was altered in 32% of precursor B cell acute lymphocytic leukemia (ALL) cases (13). In Ph-positive-ALL patients, an additional 9p abnormality [including i(9q)(q10)] had a negative impact on disease-free survival (14-16). This predicts a poor prognosis for the patient discussed in the present study; however, extended follow-up is required to confirm this.

Due to patient being <40 years old, they were subjected to a pediatric Ph-positive-ALL treatment protocol, as recommended (17). He received imatinib (400 mg daily) and 4 drugs as induction therapy: Daunorubicin (25 mg/m² weekly for 4 doses), vincristine [2 mg/week intravenously for 4 doses], prednisone (60 mg/m² for 28 days) and asparaginase (2,500 U/m² on day 6). He also received intrathecal central nervous system prophylaxis chemotherapy with cytarabine on day 1 and methotrexate on days 8 and 29. A complete remission was achieved on day 29, which was confirmed by flow cytometry, cytogenetic and molecular testing results, with minimal residual disease (<0.1%) and undetectable levels of BCR-ABL1 mRNA. The patient then continued to receive 400 mg imatinib daily. Consolodation therapy commenced on day 36 of induction. Phase one consisted of etoposide (100 mg/m²) and ifosfamide (1,800 mg/m²), daily for 5 days. Intrathecal therapy was administered on days 8 and 15 of consolidation therapy, and consisted of methotrexate (15 mg), hydrocortisone (15 mg) and cytarabine (30 mg). Phase two of consolidation consisted of high-dose methotrexate (5,000 mg/m²) on day 1 and high-dose cytarabine (3,000 mg/m² every 12 h on days 2 and 3 (4 doses in total). Triple intrathecal therapy was administered on day 1. The patient has recently undergone tissue-matched stem cell transplantation.

In conclusion, the present study reports a case of TKI-resistant Ph-positive-CML presenting with lymphoblastic crisis wherein the blast cells, in addition to the Ph chromosome, exhibited additional novel combined cytogenetic abnormalities. This report adds to the literature on newly identified TKI-resistance-conferring cytogenetic abnormalities, and also links them to precursor B cell lymphoblastic crisis. This also has its own therapeutic implications since the blast phenotype determines the treatment protocol.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

HAK wrote the manuscript, drew the figures, designed the study, interpreted the data, contributed to addressing all questions related to the accuracy and integrity of this study, and initiated this research project and interpreted the cytogenetic results. AA followed up the patient and provided the test results. All authors read and approved the final manuscript.

Ethics approval

Approval from Johns Hopkins Aramco Healthcare (JHAH) Institutional Review Board and Ethics Committee was obtained to publish this case report.

Patient consent for publication

Written informed consent was obtained from the patient for publication of this case report and any accompanying figures.

Competing interests

The authors declare that they have no competing interests.

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