RESEARCH ARTICLE

JAK-2 V617F Mutational Analysis in Primary Idiopathic Myelofibrosis: Experience from Southern Pakistan

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Abstract

Background: Primary idiopathic myelofibrosis (PMF) is a clonal Ph-chromosome negative myeloproliferative neoplasm characterized by dysregulated kinase signaling and release of abnormal cytokines. In the recent past, following JAK2 V617F mutation invention, important revolution has been made in the molecular diagnostic biology of this disease. The rational of this study was to determine the mutational status of JAK2 V617F in Pakistan patients with PMF. Materials and Methods: In this cross sectional study, 20 patients with PMF were enrolled from January 2011 to December 2014. Diagnosis was based on WHO criteria for PMF. All patients were screened for G-T point mutation (V617F) in the JAK2 gene on chromosome 9 by allele specific PCR. Results: The mean age was 57.9 ± 16.5years. The male to female ratio was 3:1. The frequency of JAK2 V617F positivity in our PMF patients was found to be 55%. Positive correlations of JAK2 V617F mutation were established with high TLC count, raised LDH and marked splenomegaly (P<0.05). No correlation of JAK2 V617F could be established with age and gender (P>0.05). Conclusions: The JAK2 V617F mutation frequency in our PMF patients was similar to those reported previously. In our hands JAK2 V617F mutated patients expressed an aggressive disease phenotype. Screening for the mutation in all suspected PMF cases could be beneficial in differentiating patients with reactive and clonal marrow fibrosis.

Keywords: Primary idiopathic myelofibrosis - JAK-2 V617F - mutational analysis - Pakistan

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Introduction

Myeloproliferative neoplasms (MPN) are a group of hematological neoplasms that share similar molecular, cellular and phenotypic abnormalities (Sag et al., 2015; Yang et al., 2015). Primary idiopathic myelofibrosis is a heterogeneous disease, characterized by dysregulated kinase signaling and release of abnormal cytokines resulting in debilitating constitutional symptoms, poor quality of life and high degree of morbidity.

The anticipated incidence of PMF is 0.5-1.5 per 100,000 individuals (Mesa et al., 1999; Tefferi., 2000). It occurs most commonly in the sixth to seventh decade of life and both genders are nearly equally affected (Thiele et al., 2008). The median age at diagnosis is around 66 years (Mitra et al., 2013).

In 2005, four research groups identified a novel single acquired mutation in the JAK2 gene on chromosome 9 which has been shown to be associated with a wide spectrum of chronic myeloproliferative neoplasm (Baxter et al., 2005; James et al., 2005).

This newly identified acquired somatic point mutation is a G-C to T-A transversion, resulting in the substitution of valine by phenylalanine at codon 617 (JAK2V617F) (Sazawal et al., 2010). JAK2 plays a role in downstream signaling pathways, such as the signal transducer and activator of transcription (STAT) pathway that is involved in cytokine signaling pathway.

JAK2V617F has been recognized in subsets of each Ph-chromosome negative MPNs. Different studies reported a variable prevalence of JAK2 V617F mutation ranging from 35-57% for PMF (Poopak et al., 2013). JAK2 V617F mutation contributed to high hemoglobin, higher white blood cell count, larger spleen span and greater need for cytoreductive treatment (Barosi et al., 2007; Poopak et al., 2013; Liu et al., 2015).

Since, there is no study available from southern Pakistan; we sought to look for the prevalence of JAK2V617F mutation in Pakistani patients with primary myelofibrosis and also to determine its correlation with age, gender, hematological and biochemical parameters.

Materials and Methods

This descriptive cross sectional study, extended from January 2011 to December 2014. Twenty patients with primary idiopathic myelofibrosis were enrolled in the present study. An informed consent was obtained from all the participating patients.

Patients were diagnosed to have PMF according to the
World Health Organization (WHO) criteria (Thiele et al., 2008). Diagnosis requires meeting all 3 major criteria and any of 2 minor criteria.

Major criteria includes

i) Presence of megakaryocyte proliferation and atypia, accompanied by fibrosis, or in the absence of fibrosis, the megakaryocyte changes must be accompanied by an increased bone marrow cellularity. ii) Not meeting WHO criteria for other myeloid neoplasm. iii) Demonstration of JAK2 V617F or in its absence no evidence of bone marrow fibrosis due to underlying inflammatory or other diseases.

Minor criteria

i) Leukerythroblastic blood picture. ii) Raised serum lactate dehydrogenase level. iii) Anemia. iv) Splenomegaly.

Complete blood counts were determined by automated hematology analyzer Cell Dyne Ruby (Abott, Diagnostics). All peripheral blood smears were reviewed by specialist hematopathologists. Biochemical test including serum creatinine, lactate dehydrogenase (LDH) and serum uric acid were detected by Hitachi 912 (Japan) by photometric assay. JAK2 V617F mutational analyses were done by Polymerase chain reaction (PCR). Bone marrow aspirate and trephine biopsy specimen were taken with Jamshidi needle.

The ethical endorsement of the study was granted by research and ethical committee LNH taken prior to the study.

Data analysis

The demographic data, clinical characteristics, and laboratory results were analyzed by descriptive analysis. Data was compiled and analyzed using SPSS version 22. The results are expressed as mean±SD for quantitative variables and qualitative variables are presented as frequency & percentages. Student ‘t’ test was applied for the comparison of mean. Data were considered statistically significant at P value < 0.05. Chi-square test was applied to assess the correlation.

Results

A total of 20 confirmed Primary idiopathic myelofibrosis patients using the non probability consecutive sampling were included in this study.

Out of 20 patients, 15 were males (75%) and 5 were females (25%) with male to female ratio of 3:1. The mean age was 57.90±16.51 years (range 22-87) years with the median age of 60 years. Mostly patients (90%) were symptomatic and presented with constitutional symptoms. In symptomatic patients, major complaints were weakness (80%); weight lost (75%); abdominal discomfort (60%); night sweats (13%); cardiovascular accident (5%) and pruritus (5%).

Physical examination revealed splenomegaly as predominant finding detected in 17 patients (85%) with the mean splenic span of 22.17±2.04cm. The hepatomegaly was detected in 11 (55%) patients followed by pallor in 7 (35%) patients.

Discussion

Primary myelofibrosis is an uncommon hematopoietic malignancy with poorest prognosis of all myeloproliferative neoplasms (Martí-Carvajal et al., 2015). A distinctive feature of PMF is dysregulation of the Janus kinase (JAK)/signal transducer and activator of transcription signaling pathway, particularly associated with the JAK2 V617F mutation, accountable in around 50% of patients with PMF (Stein et al., 2014).

In the present study, JAK2 V617F mutation was evaluated in Pakistani patients with PMF. It was noted that the prevalence of JAK2V617F mutation in our patients from southern Pakistan was analogous with that was reported in the previous local study (50%) from Northern Pakistan (Sadiq et al., 2013).

JAK2 V617F positivity in PMF has been observed from various ethnic backgrounds ranging from as low as 15% to as high as 76% (Poopak et al., 2013; Arana et
When compared with earlier reports, our results are in concurrence with regional studies reported from India; 58.8% and 52% for JAK2V617F mutational expression (Sazawal et al., 2010; Singh et al., 2015).

Other studies from Egypt and Romania by Ayad and Tevet et al have reported the frequency of JAK-2 positivity in 46% and 53.4% of patients with PMF respectively (Ayad and Nafea., 2011; Tevet et al., 2015). Another recent study from Thailand interestingly reported 100% positivity for JAK-2 mutation in Thai patients (Duangnapasatit et al., 2015). This high prevalence could be attributed to very small number (6 patients) of patients in their series (Duangnapasatit et al., 2015). However Zhang et al from China disclosed relatively low frequency for JAK-2 was detected in 40% of Chinese patients with PMF (Zhang et al., 2015).

Compared with data from developed countries outcome are more or less similar. Recent study from Taiwan also reported the similar prevalence (50%) in their patients that is comparable to our findings (Chen et al., 2015). Subsequently Cross from United Kingdom also reported mutational frequency as 60% in a large cohort of patients (Cross, 2011). Mutation frequency was determined as 54% for JAK2V617F in recent large series from the Italian group (Tefferi et al., 2014).

JAK2 mutational expression clearly segregates the disease spectrum into two broad categories. It has been also reported previously that PMF cases with JAK-2 positivity displays a higher leucocyte count and haematocrit value as compared with negative phenotype. These concerns suggest that JAK-2 expression reflect a JAK2 positivity displays a higher leucocyte count and haematocrit value as compared with negative phenotype. These concerns suggest that JAK-2 expression reflect a JAK-2 mutation with high lactate dehydrogenase which is in concurrence to our results (Larsen et al., 2007).

In conclusion, our findings are analogous to studies reported from various part of world. Marked splenomegaly, high TLC and raised LDH levels indicate that our patients had clinically advanced disease. More so is JAK2 V617F mutated patients pointing an aggressive disease phenotype. We support the recommendation that mutational screening for JAK2V617F should be incorporated into the initial evaluation of patients with suspected PMF.

References


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