Significance Of Microcytosis In The Clinical Course Of Erythrocytosis
H Fujita, S Kurosawa, S Nishimura, J Tomiyama, T Hamaki, A Ohwada

Citation

Abstract
Aim: Polycythemia vera (PV) usually occurs in people aged 60–79 years. It is difficult to diagnose PV among patients with erythrocyosis, because the test of JAK2 V617F mutation was not regularly performed in general or geriatric hospitals. Therefore, we noticed microcytosis in PV and retrospectively studied the clinical course of microcytosis in patients with erythrocytosis. Methods and subjects: The medical data of 19 patients with erythrocytosis (Hb: males, >18.5 g/dL; females, >16.5 g/dL) collected between August 2005 and December 2009 were retrospectively reviewed for parameters such as age, gender, laboratory test results (hematology, serum ferritin, and the JAK2 V617F mutation), and the day of microcytosis onset during a 2-year period. The patients were divided into 2 groups depending on the presence or absence of the JAK2 V617F mutation, and the day of microcytosis onset during a 2-year period. The patients were divided into 2 groups depending on the presence or absence of the JAK2 V617F mutation. Microcytosis was defined as low mean corpuscular volume (MCV; <79 fL). Results: MCV was significantly lower in the JAK2 V617F -positive (PV, N=10) than in the negative erythrocytosis group (N=9). Microcytosis was noted in 2 among PV patients at initial presentation, whereas all patients exhibited microcytosis within 2 years. On the other hand, microcytosis was not noted at initial presentation in the JAK2 V617F-negative erythrocytosis group, and only 1 patient exhibited microcytosis within 2 years because of gastrointestinal bleeding. Microcytosis in the clinical course was associated with the presence of the JAK2 V617F mutation (P < 0.05). Conclusion: Microcytosis may be a characteristic of the clinical course of PV and a good marker of the presence of the JAK2 V617F mutation.

INTRODUCTION
Polycythemia vera (PV) occurs in all age groups, although the incidence increases with age. Berlin NI reported the median age at diagnosis to be 60 years1, while Anfa B et al found that the highest incidence was in people aged 70–79 years2. Because JAK2 V617F mutations are noted in approximately 95% of the patients with PV, testing for this mutation is important for the diagnosis of this disease3. However, the test is not regularly performed in general or geriatric hospitals in several countries including Japan4. Before publication of the World Health Organization (WHO) criteria in 2008 that included the presence of the JAK2 V617F mutation5, definite diagnosis of PV was based on various guidelines6-7. Splenomegaly and the number of leukocytes and platelets are very useful markers in the initial clinical diagnosis of PV6. PV is occasionally accompanied by iron deficiency, which results in microcytosis (i.e., low mean corpuscular volume [MCV])8. Therefore, we investigated the occurrence of microcytosis from the perspective of initial diagnosis of PV.

Phlebotomy is used as an initial treatment for PV9. This procedure was performed for unclassified polycythemia before the development of tests for the JAK2 V617F mutation, serum erythropoietin level, etc. Venesection alone is known to cause iron deficiency in cases of PV9. However, patients with this blood disorder may or may not exhibit hematological microcytosis. Microcytosis in adulthood is known to occur in conjunction with anemia because of iron deficiency, chronic diseases, hemoglobinopathy, etc.11,12 Mutations in the ank/ank gene induce microcytosis that is unrelated to the JAK2 V617F

1 of 6
Significance Of Microcytosis In The Clinical Course Of Erythrocytosis

Moreover, the detailed clinical course of microcytosis in patients with PV remains unclear. Therefore, we retrospectively studied the clinical course of microcytosis in patients with PV. Microcytosis occurs more frequently during the clinical course of PV than during that of JAK2 V617F-negative erythrocytosis. First, we determined the significance of microcytosis in the clinical course of PV. Microcytosis during the clinical course of PV may be a result of iron deficiency or excessive erythropoiesis due to the JAK2 V617F mutation. Furthermore, it may be a good marker of the presence of the JAK2 V617F mutation.

METHODS AND SUBJECTS

CLINICAL EVALUATION

Tokyo Metropolitan Bokutoh Hospital is a 729-bed acute-care hospital located in eastern Tokyo. We retrospectively reviewed the medical data of 19 patients with erythrocytosis (Hb: males, >18.5 g/dL; females, >16.5 g/dL) collected between August 2005 and December 2009. The patients were divided into 2 groups according to the presence or absence of the JAK2 V617F mutation. Microcytosis was defined as low mean corpuscular volume (MCV) <79 fL. The patients' charts were reviewed for age, gender, gastrointestinal bleeding events, laboratory test results (i.e., hematology, serum ferritin level, and the JAK2 V617F mutation), and the number of phlebotomies received during 2 years. The presence of the JAK2 V617F mutation was determined from venous blood from patients with erythrocytosis by the SRL company (Tokyo, Japan). The diagnostic criteria for PV were based on the revised WHO classification (1). All JAK2 V617F mutation-positive patients (n = 10) satisfied the WHO criteria. On the other hand, the mutation-negative patients (n = 9) could not be diagnosed with PV despite high hemoglobin concentrations (serum erythropoietin:15±2 mIU/mL). Patients with secondary polycythemia due to chronic obstructive pulmonary or cyanotic heart diseases or with high serum erythropoietin (>29 mIU/mL) were excluded from this study. Phlebotomies were performed in our hospital if patients had high hematocrit levels (>55%) and were repeatedly performed until hematocrit levels were <45%. The amount of phlebotomized whole blood was 400 mL per patient per phlebotomy. Hydroxycarbamide was administered for phlebotomy-resistant elderly patients (>60 years old) or high-risk groups including those with previous thrombosis. However, all 19 patients had neither previous thrombosis nor episodes of vascular events over the past 2 years. All study participants provided informed consent, and the study design was approved by the ethics review board of our institution.

STATISTICAL ANALYSIS

We compared differences between groups using Wilcoxon analysis. The data represent the mean ± standard error (SE). The transient incidence of microcytosis in the clinical course between the groups was determined by Chi squared analysis (log rank). All statistical tests were conducted using JMP version 8.0 (SAS Institute, Inc., Cary, NC, USA). The level of statistical significance was set at P < 0.05.

RESULTS

Clinical features of PV or JAK2 V617F-negative erythrocytosis

Among the 19 patients with erythrocytosis, 10 were positive for the JAK2 V617F mutation and were diagnosed with PV by meeting the WHO criteria. However, 9 patients with high hemoglobin concentrations tested negative for the JAK2 V617F mutation and were not diagnosed with PV. The patients with JAK2 V617F-positive erythrocytosis, namely PV (age, 66 ± 3 years), were significantly older than those without the JAK2 V617F mutation (56 ± 3 years); however, no gender difference was noted between groups (male/female: PV, 6/4; JAK2 V617F-negative erythrocytosis, 8/1) (Table 1). None of the patients with PV had gastrointestinal bleeding during the 2-year period; only 1 patient who did not have the mutation experienced this.

In the laboratory tests performed at initial presentation, the leukocyte, erythrocyte, and platelet counts were significantly higher in the PV than in the JAK2 V617F-negative erythrocytosis (Table 1). Hemoglobin concentration and hematocrit were similar between groups (Table 1).

Clinical course of microcytosis in PV and JAK2 V617F-negative erythrocytosis

Among the 19 patients with erythrocytosis, only 2 with PV (2/10, 20%) exhibited microcytosis at the initial presentation, whereas none among the mutation-negative erythrocytosis did (0/9, 0%). Moreover, MCV was significantly lower in the PV than in the mutation-negative erythrocytosis (Table 1). The level of serum ferritin at the initial presentation was significantly lower in the PV (27.0 ± 4.8 µg/L, n=8) than in the mutation-negative erythrocytosis (126.7± 26.8 µg/L, n=7), although we only examined serum ferritin levels in 15 among 19 subjects at the initial presentation.

Within the 2 year period, 8 patients with normocytic PV at
the initial presentation exhibited transient microcytosis (8/8; 100%) throughout the initial presentation to days ~23–492 (Figure 1); However, only 1 mutation-negative patient exhibited this (1/9, 11%) at day 265.

Regarding the hematology of the microcytosis patients, erythrocyte count, hemoglobin concentration, and hematocrit (RBC: 6780 ± 190 × 10^9/L, Hb: 16.7 ± 0.6 g/dL, Hct: 52 ± 2%) were significantly lower than those measured at the initial presentation. However, the leukocyte (13.2 ± 0.2 × 10^9/L) and platelet (500 ± 100 × 10^9/L) counts were not significantly different from those at the initial presentation. Serum ferritin levels significantly decreased in the 8 patients with microcytosis (17.0 ± 2.9 µg/L).

Phlebotomies were similarly required in both the PV (10/10, 7 ± 2 times) and mutation-negative erythrocytosis (9/9, 6 ± 2 times) during the 2-year period.

Hydroxycarbamide administration was required by 5 out of the 10 patients with PV (50%). After hydroxycarbamide therapy, 4 of 5 patients (80%) exhibited normocytosis ranging from days 73–221. Serum ferritin levels increased in the 5 patients with normocytosis (50.0 ± 8.7 µg/L). Two of the four patients who exhibited normocytosis again exhibited microcytosis at the initial presentation. Six of 10 patients with PV still had microcytosis during the 2-year period. On the other hand, 1 out of the 9 patients with JAK2 V617F mutation-negative erythrocytosis exhibited microcytosis on day 265, presented with normocytosis again on day 724 because of the resolution of gastrointestinal bleeding without iron supplementation. Therefore, all JAK2 V617F mutation-negative erythrocytosis patients finally exhibited normocytosis by the end of the 2-year period.

Microcytosis among the patients with erythrocytosis at the initial presentation was not associated with the presence of the JAK2 V617F mutation (P = 0.26). However, transient microcytosis in the clinical course was significantly associated with the presence of the JAK2 V617F mutation (P < 0.05).

### DISCUSSION

#### CLINICAL FEATURES OF PV

Among the patients of the PV group that exhibited microcytic characteristics (2 and 8 cases during the initial presentation and clinical course respectively), red blood cell, leukocyte, and platelet counts were greater in comparison with those in the JAK2 V617F mutation-negative erythrocytosis group (9 cases, no microcytosis at the initial presentation). There was no significant difference in the
Significance Of Microcytosis In The Clinical Course Of Erythrocytosis

Microcytosis in the clinical course of PV may be a result of iron deficiency and excessive erythropoiesis due to the JAK2 V617F mutation. In addition, microcytosis in erythrocytosis may be a good marker for the presence of the JAK2 V617F mutation. 

CONCLUSION

References
Significance Of Microcytosis In The Clinical Course Of Erythrocytosis

Author Information
Hiroshi Fujita
Department of Transfusion Medicine, Tokyo Metropolitan Bokutoh Hospital

Saiko Kurosawa
Department of Transfusion Medicine, Tokyo Metropolitan Bokutoh Hospital

Shigeko Nishimura
Department of Transfusion Medicine, Tokyo Metropolitan Bokutoh Hospital

Junji Tomiyama
Department of Internal Medicine, Tokyo Metropolitan Bokutoh Hospital

Tamae Hamaki
Department of Internal Medicine, Tokyo Metropolitan Bokutoh Hospital

Akira Ohwada
Department of Internal Medicine, Tokyo Metropolitan Bokutoh Hospital