Beyond Hemoglobin: When and How to Work Up Possible Polycythemia Vera

Gene Shaw, MD and Richard Berg, MS

Background: World Health Organization (WHO) 2017 diagnostic criteria for hemoglobin levels in polycythemia vera (PV) were lowered from 185 g/L to 165 g/L for men and from 165 g/L to 160 g/L for women, but these cutoffs were not designed for screening.

Objectives: The primary aim of this study was to assess the value of laboratory and clinical parameters in deciding whether to further pursue a diagnosis of PV. A secondary aim was to explore the diagnostic utility of bone marrow morphology.

Methods: We evaluated clinical and laboratory parameters that may be useful when considering further diagnostic work-up, emphasizing PV vs. secondary erythrocytosis (SE). We classified 200 patients with JAK2 V617F testing using WHO criteria.

Results: Patients with myeloproliferative neoplasms (MPN) were rarely under age 40 and uncommonly obese (BMI $\geq 30$ kg/m$^2$). Current smoking history favored SE, and these patients rarely had a platelet count $\geq 450 \times 10^3/\mu$L. Laboratory parameters suggesting greater PV likelihood were: RBC $> 6.8 \times 10^6$ for men or $> 5.9 \times 10^6$ for women; low erythropoietin; and low MCV or low ferritin. Bone marrow morphology (available in 111 cases) was generally more cellular in PV vs. SE and assessed disease progression.

Conclusions: Readily accessible clinical and laboratory data can assist in considering a PV workup, and a possible diagnostic algorithm is presented. These preliminary findings warrant larger studies to develop a more formal PV-risk scoring system with optimal cutoffs and weighting.

Keywords: Polycythemia vera; Secondary erythrocytosis; Myeloproliferative neoplasms; JAK2 mutations; Erythropoietin

The World Health Organization (WHO) diagnostic criteria for myeloproliferative neoplasms (MPN), including polycythemia vera (PV), were formally revised in 2017. To distinguish “masked” PV from secondary erythrocytosis (SE), the major criterion for hemoglobin (HB) level in men was significantly decreased from 185 g/L to 165 g/L, while the HB level for women was slightly lowered from 165 g/L to 160 g/L. Janus kinase 2 (JAK2) mutations (either JAK2 V617F in exon 14 or rarely mutations in exon 12) are present in virtually all PV patients. These mutations were originally included in the 2008 WHO criteria and have been retained as one of three major criteria. Bone marrow (BM) morphology has been elevated from a minor to a major criterion that emphasizes hypercellularity and pleomorphic megakaryocytes. Low serum erythropoietin (EPO) level is now the sole minor criterion.

However, in clinical practice the diagnostic process is sequential, usually with abnormalities first noted in the complete blood count (CBC), which are then correlated with the clinical history and physical exam. The new WHO HB criteria were not designed for general screening and are clearly too low for that purpose. The red blood cell count (RBC) may be underutilized. In addition, iron deficiency
may raise suspicion for PV,\textsuperscript{10} and the role of EPO warrants further investigation.\textsuperscript{11,12} Key clinical parameters include smoking history and body mass index (BMI); the latter is often associated with obstructive sleep apnea (OSA). If there is sufficient clinical suspicion for PV, JAK2 molecular testing is generally the next diagnostic step.

The primary aim of the current study was to assess the value of readily available laboratory and clinical parameters in making an informed decision on whether to further pursue a diagnosis of PV. A secondary aim was to explore the diagnostic utility of BM morphology.

**Materials and Methods**
Following Institutional Review Board approval, 200 cases of suspected MPN, other than chronic myeloid leukemia, that occurred after 2008 (when JAK2 testing became widely available) in the Marshfield Clinic system were identified. The following inclusion criteria were used: age 18 years or older, a JAK2 V617F mutation test result, and available clinical follow-up information. An exclusion criterion was ≥ 5% blasts in the BM at the time of MPN diagnostic evaluation.

The white blood cell (WBC) count at the time of initial diagnostic evaluation was recorded along with the highest values on or prior to that date for Hb, RBC, and platelets along with the lowest values for MCV, EPO, and ferritin. Transiently abnormal values due to a clinical condition (eg, blood loss) were not included.

For BM evaluation, the following data were obtained from the medical record: cellularity, myeloid:erythroid (M:E) ratio, megakaryocyte morphology (increased and/or abnormal), presence or absence of stainable iron, reticulin fibrosis (Grades MF-0, MF-1+, MF-2-3+), and cytogenetics. The following clinical data were gathered: age, gender, smoking history (never, former, or current), splenomegaly (Y/N), BMI (kg/m\(^2\)), history of OSA (Y/N), treatment with phlebotomy (Y/N), and survival (alive or date of death).

Two hematopathologists independently reviewed the bone marrow without knowledge of patient age or category assignment and recorded the following information: evaluable biopsy length, cellularity in 10% increments (if biopsy inadequate, clot sections used if adequate), megakaryocyte morphology (increased: Y/N; abnormal: Y/N; predominant type of abnormality: staghorn, cloud-like, or unclear), and reticulin fibrosis (WHO MF- 0, 1, or 2-3).

The following scheme for hypercellularity was adopted: age 18-39 years, ≥ 70%; 40-69 years, ≥ 60%; and age ≥ 70 years, ≥ 50\%\textsuperscript{13,14} The EPO method used in our laboratory was slightly modified in December 2013. Prior to that, the reference range was 4-24 mIU/mL; subsequently it was 3-19 mIU/mL.

Mayo Clinic Laboratories performed the vast majority of JAK2 V617F testing for this study and reported a qualitative result prior to November 2011. Other reference laboratories gave a qualitative result throughout the study. Quantitative results were reported with an equivocal range of 0.06%-0.6%.

Applying 2017 WHO criteria and subsequent clinical course, patients were placed into one of seven categories: PV, post-PV

![Figure 1. Patient BMI versus diagnostic category (Grey diamond-shaped areas encompass quartiles 2 and 3).](image-url)
Table 1 Patient characteristics and laboratory data by diagnostic group

|                        | PV  | Post-PV | MF         | SE        | ET        | Overt | PMF | PreMF  | Other |
|------------------------|-----|---------|------------|-----------|-----------|-------|-----|--------|-------|
| Total Patients         | 36  | 7       | 85         | 25        | 16        | 12    | 19  |
| Male (%)               | 55.6| 57.1    | 60.0       | 36.0      | 68.8      | 58.3  | 63.2|
| Median age (yrs)       | 73.3| 66.7    | 57.5       | 73.5      | 72.5      | 79.9  | 56.4|
| Minimum                | 23.7| 59.4    | 16.3       | 22.8      | 44.6      | 50.2  | 26.5|
| Maximum                | 88.6| 86.3    | 86.2       | 92.3      | 90.3      | 97.1  | 82.1|
| Median BMI (kg/m²)     | 27.5| 30.8    | 30.8       | 26.3      | 27.0      | 27.7  | 31.4|
| Minimum                | 17.2| 25.1    | 19.3       | 17.1      | 17.6      | 21.7  | 20.4|
| Maximum                | 35.8| 37.1    | 53.5       | 39.7      | 44.8      | 40.2  | 46.8|
| Any smoking (%)        | 58.3| 28.6    | 69.4       | 36.0      | 68.8      | 66.7  | 68.4|
| JAK2+ (%)              | 100 | 100     | 0.0        | 64.0      | 62.5      | 58.3  | 31.5|

Hb (g/L) Females

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 172    | 142     | 206     |
| Male                   | 153    | 139     | 183     |
| Female                 | 176    | 129     | 203     |
| Minimum                | 142    | 111     | 151     |
| Maximum                | 159    | 159     | 159     |

Hb (g/L) Males

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 181    | 142     | 228     |
| Male                   | 191    | 170     | 242     |
| Female                 | 187    | 156     | 221     |
| Minimum                | 142    | 102     | 173     |
| Maximum                | 161    | 159     | 206     |

RBC (x10⁶/µL) Females

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 6.0    | 4.7     | 10.3    |
| Male                   | 5.3    | 4.4     | 7.5     |
| Female                 | 5.6    | 5.0     | 7.9     |
| Minimum                | 4.7    | 3.3     | 5.7     |
| Maximum                | 4.8    | 3.8     | 5.9     |

RBC (x10⁶/µL) Males

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 6.9    | 5.5     | 10.3    |
| Male                   | 6.8    | 5.3     | 7.5     |
| Female                 | 6.0    | 5.2     | 7.9     |
| Minimum                | 5.5    | 3.0     | 5.7     |
| Maximum                | 5.4    | 4.5     | 5.9     |

Platelets (x10³/µL)

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 686.0  | 164.0   | 1598.0  |
| Male                   | 617.0  | 145.0   | 1501.0  |
| Female                 | 283.0  | 121.0   | 757.0   |
| Minimum                | 164.0  | 123.0   | 2437.0  |
| Maximum                | 864.5  | 247.0   | 1747.0  |

MCV (fl)

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 84.2   | 55.6    | 94.5    |
| Male                   | 76.2   | 65.8    | 86.6    |
| Female                 | 89.1   | 64.3    | 102.0   |
| Minimum                | 84.7   | 71.5    | 97.4    |
| Maximum                | 87.8   | 82.4    | 95.7    |

EPO (mIU/mL)

|                        | Median | Minimum | Maximum |
|------------------------|--------|---------|---------|
| Total                  | 2.5    | 0.6     | 14.0    |
| Male                   | 4.0    | 1.5     | 4.0     |
| Female                 | 7.3    | 1.0     | 8.0     |
| Minimum                | 13.3   | 5.0     | 100.0   |
| Maximum                | 18.0   | 18.0    | 18.0    |

BMI, body mass index; Hb, hemoglobin; EPO, erythropoietin; MCV, mean corpuscular volume; RBC, red blood cell; PV, polycythemia vera; MF, myelofibrosis; SE, secondary erythrocytosis; ET, essential thrombocythemia; PMF, primary myelofibrosis; PrePMF, prefibrotic primary myelofibrosis
myelofibrosis (post-PV MF), SE, essential thrombocythemia (ET), prefibrotic primary myelofibrosis (prePMF), overt PMF, or “other”. The JAK2 V617F mutational burden was not used in category assignment.

Statistical comparisons were based on standard methods, including chi-square tests for categorical outcomes (eg, smoking status) and the Kruskal-Wallis test for continuous outcomes (eg, lab results). Since we did not have specific a priori hypotheses, and since we had no gold-standard disease classification, the results will require confirmation through additional studies.

Results

For all patients in the Marshfield Clinic system who had a Hb level measured during 2016, the first value was > 165 g/L in 7.7% of 62,000 men and > 160 g/L in 1.4% of 84,000 women. In contrast, using the previous, more stringent 2008 WHO criteria, only 0.3% of men had Hb > 185 g/L, while 0.6% of women had Hb > 165 g/L.

Table 1 summarizes the clinical and laboratory data for the 200 patients in this study. The seven patients classified as post-PV MF were similar to the other PV patients; thus, they were combined for statistical analyses. Patients with SE, as expected, had a lower mean age. The only category with more females than males was ET. Of SE patients, 45 of 85 (53%) were current smokers versus 6 of 43 PV/post-PV MF patients (14%, $P < 0.001$). Only 3 of 96 MPN patients were under 40 years-of-age.

Of SE patients, 46 (54%) were obese (BMI > 30 kg/m²) compared with 10 (23%) of 43 PV/post-PV MF patients ($P = 0.001$) as shown in Figure 1. Sixteen of 19 SE patients with a history of OSA were obese (BMI > 30 kg/m²). Regarding obesity and cigarette smoking (either former or current) as risk factors for SE, 79 of 85 (93%) SE patients had at least one risk factor, compared with 27 of 43 (63%) PV/post-PV MF patients ($P < 0.001$).

Hb level was not helpful in distinguishing PV/post-PV MF from SE. Of the 43 PV/post-PV MF patients, 25 met the 2008 WHO Hb criteria. Another 11 men met only the 2017 criteria; 8 had a BM evaluation, and all were hypercellular. Three did not have a BM evaluation, but their further clinical course supported PV as the correct assignment.

Seven additional patients (one male and six females) were classified as PV/post-PV MF despite all prior available Hb levels being ≤ 165 g/L or 160 g/L, respectively. All seven had an elevated RBC and evidence of iron deficiency, with further clinical course supporting PV. Of 53 patients with ET or PMF, 8 (all men) had Hb levels above the 2017 WHO PV criterion, but this was felt to be SE superimposed upon their primary MPN.

Figure 2 shows RBC versus diagnostic group. Of 24 male PV/post-PV MF patients, 14 had an RBC count > 6.8 x 10⁶/L versus only 3 of 51 SE patients ($P < 0.001$). For females, 11 of 19 PV/post-PV MF patients had an RBC count > 5.9 x 10⁶/L versus 3 of 33 SE patients ($P < 0.001$).
Figure 3 shows platelet count versus diagnostic group. SE patients rarely had platelet counts $\geq 450,000/\mu L$ vs. patients with MPN ($P < 0.001$).

The MCV and ferritin levels trended lower in the PV/post-PV MF patients, but with overlap. The MCV/RBC ratio for each patient in the various categories did not show better group separation.

As shown in Figure 4, EPO levels were usually lower in PV/post-PV MF patients vs. SE, but with overlap. Of 27 PV/post-PV MF patients, 20 had low EPO levels versus 15 of 72 SE patients ($P < 0.001$). Levels above 9 mIU/mL were common in SE patients, but were seen in just one PV patient when the Hb level was 144 g/L from blood loss.

All 43 PV/post-PV MF patients were JAK2+, with just one having an exon 12 mutation. He had a classic PV clinical presentation and clinical course. For ET, prePMF, and overt PMF, a slight majority (16/25, 10/16, and 7/12, respectively) were JAK2 V617F+.

Among 19 patients classified as “other” were four patients with myelodysplastic/myeloproliferative neoplasm with ring sideroblasts and thrombocytosis (MDS/MPN-RS-T). Another four had an unclassifiable MDS or MPN. One patient had congenital erythrocytosis. Another had a presentation like PV with low EPO, but was negative for JAK2 V617F and exon 12 mutations. Another patient was favored to have SE, but had equivocal JAK2 V617F results. Lastly, eight patients likely had a reactive cause for their elevated counts.

Eight patients had JAK2 V617F values in the equivocal range, but follow-up testing did not show an upward trend, and none were diagnosed with an MPN. Five were classified as SE based on other laboratory data and clinical presentation. One PV patient initially had a false negative JAK2 V617F test by an allele-specific assay. Subsequent sequencing actually found the V617F mutation along with a second mutation in exon 14 that likely disrupted primer binding.

Cytogenetic testing was performed on 112 patients with an available BM aspirate, and 17 had cytogenetic abnormalities associated with a myeloid neoplasm. A 20q deletion was the most common abnormality, seen in seven cases. Eight of 21 patients with MPN having myelofibrosis (combining post-PV MF and overt PMF) had abnormal cytogenetics.

Three PV/post-PV MF patients went on to develop acute myeloid leukemia, and one developed a high grade myeloid neoplasm with 14% BM blasts. One overt PMF patient (“triple negative” for JAK2V617F, CALR, and MPL) developed acute myeloid leukemia.

Of nine JAK2 V617F-negative ET patients, eight were tested for a CALR mutation, and two were positive. One ET patient had a myeloproliferative leukemia (MPL) Trp515Leu mutation. Of 11 prePMF or overt PMF patients that were

![Figure 3](image-url)  
*Figure 3. Highest previous platelet count versus diagnostic category (Grey diamond-shaped areas encompass quartiles 2 and 3).*
JAK2 V617F-negative, only five had CALR testing, and three were mutated.

Very few patients had red cell mass measurements. One patient was erroneously considered Gaisbock syndrome (erythrocytosis putatively due to plasma volume contraction) based on a normal red cell mass. Eventually he was diagnosed with PV, confirmed by a JAK2 V617F mutation.

There were 111 BM specimens available for review. In 13 cases, there was either no biopsy or it was of inadequate length (≤ 0.2 cm). Of the remaining 98 biopsies, 61 were 0.3–0.9 cm in evaluable length, 28 were 1.0–1.4 cm, and 9 had the WHO recommended ≥ 1.5 cm.15 For reticulin fibrosis, 18 of 91 cases that could be assessed were called MF-0 or MF-1+ by one hematopathologist, but MF-2-3+ by the other, which could influence diagnostic category assignment. The subjective assessment of megakaryocytes as having “staghorn” or “cloud-like” nuclei showed poor concordance.

Of 43 patients with PV/post-PV MF, 31 had adequate BM samples to evaluate cellularity. In 28 patients, it was interpreted as hypercellular by our hematopathologists. Megakaryocytes were felt to be increased in all PV/post-PV MF patients and abnormal in all but two. Of the 85 SE patients, 26 underwent BM evaluation. BM was called hypercellular in just three patients (P < 0.001 when compared with PV/post-PV MF.

Of 16 patients with overt PMF, 15 had BM evaluation, with hypercellularity for age in 13. Four of the 16 had a WBC ≥ 11 x 10^9/µL. For prePMF, 11 of 12 had BM evaluation, with nine being hypercellular. Nine of the 12 patients had a WBC count ≥ 11 x 10^9/µL.

Of 25 ET patients, 18 had a BM evaluation; three were slightly hypercellular for age. Six of the 25 had a WBC ≥ 11 x 10^9/µL. None of 25 ET patients had splenomegaly, versus 5 of 12 prePMF patients.

Based on visual inspection of the scatter plots and statistical comparisons cited above, threshold values were chosen to assist in differentiating PV from SE. RBC > 6.8 x 10^6/µL for men or > 5.9 x 10^6/µL for women, platelets ≥ 450 x 10^3/µL, EPO < 4 mIU/mL, and MCV < 80 fl (or ferritin < 24 ng/mL) increased the probability of PV vs. SE. Age < 40 years, BMI > 30 kg/m² (and especially > 35 kg/m²), and EPO > 9 mIU/mL reduced the probability.

The following PV risk scoring system was devised as an exploratory investigation:

- Age < 40 years: -1
- RBC > 6.8 x 10^6/uL (men) or > 5.9 x 10^6/uL (women): +1
- Platelets ≥ 450 x 10^3/uL: +1
- BMI 30-35 kg/m²: -0.5
- BMI > 35 kg/m² or history of OSA: -1
- Smoking: currently or chronic obstructive pulmonary disease (COPD): -0.5
- MCV < 80 fl or ferritin < 24: +0.5
- EPO* < 4 mIU/mL: +0.5
- EPO* > 9 mIU/mL: -1

*EPO is often not available at initial patient presentation.

**Figure 4.** Erythropoietin levels versus diagnostic category. (Grey diamond-shaped areas encompass quartiles 2 and 3.)
Without EPO, a total score of $\geq 0.5$ yielded a sensitivity of 93% and specificity of 94% for diagnosing PV. Incorporating EPO levels improved both sensitivity and specificity to 98% using the same total score cutoff.

**Discussion**

The prevalence of PV in the United States was estimated at 22 in 100,000 individuals in 2008. Based on the Hb data in the Marshfield Clinic system cited above, roughly 1 in 350 males and 1 in 64 females with Hb $> 165$ g/L or $> 160$ g/L, respectively, would be expected to have PV. Thus, use of the new Hb criteria to trigger JAK2 testing, without consideration of other clinical factors, could result in excessive testing. In this study, 74% of PV patients had a low EPO level, which corresponds with a recent review. However, the specificity (when applied to SE patients) was 79%, which is considerably below the 92%–99% cited in that review. In our study, a few PV patients were identified who had high EPO levels during a time of transient blood loss, which is an appropriate physiologic response. Similarly, patients who undergo therapeutic phlebotomy will eventually raise their EPO levels provided renal function is normal. Therefore, EPO levels need to be obtained when the Hb level is at least high-normal.

The RBC count is a useful parameter in separating PV from SE. In our study, about 60% of PV patients had values $> 6.8 \times 10^6/\mu L$ for males and $> 5.9 \times 10^6/\mu L$ for females, whereas few SE patients exceeded these cutoffs (Figure 2). Although, thalassemia often has a high RBC and low MCV, in clinical practice this rarely poses a problem. Thalassemia is a genetic disorder characterized by a lifelong high RBC and low MCV without an elevated Hb level.

Many laboratories have an equivocal range for the JAK2 V617F mutation. Values in this range should be repeated at a
Whether true JAK2-negative PV exists is controversial.\textsuperscript{22,23} However, since only about 2\% of PV patients have an exon 12 mutation, it is debatable whether all JAK2 V617F-negative patients being evaluated for PV should have this as a follow-up test.

As a criterion for MPN diagnosis, BM morphology presents several challenges. Well-validated age-adjusted cutoffs for BM cellularity have not been established to diagnose MPN.\textsuperscript{13,14} The current study used a simple scheme for age-adjusted hypercellularity that could be adopted in routine practice. Reactive/regenerative changes can also increase BM cellularity. Furthermore, the biopsy may be small with crush artifact, procedure-related hemorrhage, or have hypocellular subcortical areas not apparent from the biopsy orientation.

In our study with limited morphologic evaluation, the subtyping of megakaryocytes (ie, staghorn or cloud-like) was unreliable.\textsuperscript{24} Thus, the distinction between ET and prePMF was essentially based on BM cellularity, WBC count, and splenomegaly; M:E ratio was not helpful. A recent study of 211 BM biopsies independently reviewed by two hematopathologists found low reproducibility of the WHO histologic criteria for MPN, even though all biopsies were $\geq 1.5$ cm in length.\textsuperscript{25}

Cellularity is best evaluated from the biopsy, ideally $\geq 1.5$ cm in length without crush artifact;\textsuperscript{15} however, in many practice settings biopsies are usually smaller.\textsuperscript{26} Clot sections are suboptimal, but can be used if they have moderately sized spicules $\geq 0.2$ cm. Aspirate smears can only give a qualitative estimate.

In the appropriate clinical context, an MPN “driver mutation” (ie, JAK2, CALR, or MPL) establishes the presence of a myeloid neoplasm.\textsuperscript{23} BM evaluation may yield important prognostic information; e.g. percentage of marrow blasts, dysplasia, ring sideroblasts, and marrow fibrosis. Also, BM generally provides the best sample for a cytogenetic study.

Some hematologists may be reluctant to embrace BM evaluation as a mandatory major criterion for PV diagnosis. For example, in a JAK2 V617F+ patient with an elevated Hb and no reason for SE, a diagnosis of PV is reasonably secure without BM evaluation. However, the presence of increased poikilocytosis (including nucleated RBCs or teardrop forms), a myeloid left-shift (possibly including circulating blasts), or splenomegaly would raise concern of myelofibrosis or disease progression, and bone marrow evaluation would be clearly indicated.

Conventional cytogenetics continue to play a key role in the diagnosis and prognostication of myeloid neoplasms.\textsuperscript{27,28} The finding of relatively frequent cytogenetic abnormalities in MPN patients with marrow fibrosis (overt PMF and post-PV MF) fits with the observation that these patients have greater overall genetic complexity contributing to their unfavorable prognosis.\textsuperscript{29-31}

When facing a differential diagnosis of PV vs. SE, the simple scoring system provided in this study may assist in deciding when to proceed to further diagnostic testing. Relevant clinical and laboratory information available in nearly all patients includes age, gender, RBC, platelet count, MCV, and BMI. Smoking history has limited reliability, but current smoking or a history of heavy smoking (often accompanied by COPD) is helpful. Although a history of OSA may provide additional support for SE (beyond obesity), it is often not well documented nor is its severity quantified. In contrast, BMI is more objective and increasingly documented in the medical record. Ferritin level is occasionally available and augments MCV in assessing iron deficiency.

A suggested diagnostic algorithm is presented in Figure 5 to differentiate PV from SE. A more formal PV-risk scoring system would require larger patient cohorts to perform multivariate analysis with assignment of optimal cutoffs and weighting of risk factors.

Costs of laboratory tests vary widely between individuals, depending on payer (private insurance or government) and discounts. Currently within the Marshfield Clinic Health System, JAK2V617F, JAK2 exon 12, and EPO tests have comparable fees, roughly between $150 and $200.

Our study reflects real-life clinical practice and the everyday challenges in diagnosing MPN. Limitations include patient selection that was not done in a way to provide a statistically representative sample, and patients were diagnosed at different times in the course of their disease. The small size of the subgroups limited the power of statistical comparisons. Genetic data, other than JAK2 V617F, was sporadically available. The patients were not uniformly treated, and 42\% did not have a BM evaluation. MPL and CALR tests, which became more widely available in 2009 and early 2014, respectively, were done in a relatively small number of cases.

Our patient cohort was predominantly of northern European descent from the upper Midwest of the United States at about 360 meters above sea level. Other populations may differ significantly in prevalence of obesity or cigarette smoking. Therefore, the use of these clinical parameters for differentiating PV from SE may need modification in other settings.

JAK2+ MPN may have overlapping phenotypes that evolve over time.\textsuperscript{27} In this study, several JAK2 V617F+ patients...
were difficult to classify due to the possibility of superimposed SE; a few evolved to a clear-cut PV presentation. The phenotype and prognosis is influenced by the mutational burden and additional genetic aberrations that are increasingly being elucidated by gene panel sequencing.32-36

This study was conducted using the classical paradigm for non-chronic myeloid leukemia MPN, attempting to classify each patient as either PV, post-PV MF, ET, prePMF, or overt PMF. Looking to the future, with advances in our understanding of the molecular basis of MPN, this scheme may need revision. A new model (similar to that for acute leukemias) could primarily subclassify these disorders by their driver mutation(s) and other genetic abnormalities.37

Various lines of evidence support such a change. Rather arbitrary Hb levels have been proposed to discriminate "masked" PV from JAK2+ ET.27 Yet, nearly 30% of patients presenting as JAK2+ ET progress to a PV phenotype at 15 years.38 Approximately 10% of PV and 5% of ET cases develop myelofibrosis indistinguishable from overt PMF at 15 years.39 The distinction between this evolution and overt PMF presumably evolved from prePMF is not possible if there is inadequate historical data. Moreover, such separation does not influence patient management. PMF patients with CALR mutations (mostly the more favorable type 1 with a 52 BP deletion) have a better prognosis than JAK2 V617F+ patients, who in turn do better than “triple negative” cases.39 Mutational status acquired by sequencing a panel of relevant genes may become the best predictor of future disease course and response to treatment.37,41

Conclusion
In summary, this study of 200 patients who underwent JAK2 V617F mutation testing because of a clinically suspected MPN provided several key findings, which will require confirmation by future studies.

1. The WHO 2017 Hb diagnostic criteria were not intended to be used indiscriminately to screen for PV. Hb level alone is not helpful in distinguishing PV from SE. However, PV patients often have higher RBC values (> 6.8 x 10^6/μL for men and > 5.9 x 10^6/μL for women) along with lower MCV and ferritin values than patients with SE. Platelet counts ≥ 450,000/μL are rare in SE patients.
2. The decision to “trigger” molecular testing and/or BM evaluation should incorporate clinical information. MPN patients are rarely under age 40 (only 3% in this study). The vast majority of SE patients (93% in this study) have at least one risk factor (eg, obesity [BMI ≥ 30 kg/m²] or smoking history.)
3. JAK2 V617F positivity (above each laboratory’s equivocal range) is diagnostic of MPN in the appropriate clinical context. However, equivocal values are not confirmatory and repeat testing is advised.
4. Although low EPO levels are more common in PV, this finding is not specific enough to confirm the diagnosis. Conversely, high EPO values above the reference range midpoint support SE, provided the Hb level is at least high-normal at the time of testing.
5. Currently, JAK2 exon 12 sequencing can be reserved for selected cases, rather than being a reflex order upon negative JAK2 V617F testing. This recommendation may change if costs come down.
6. BM evaluation (including a biopsy of adequate size) can assess marrow blasts, myelofibrosis, and provide a sample for cytogenetics. It may assist in distinguishing classical MPN subtypes and should be performed before therapeutic intervention.

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