Revised Myeloproliferative Neoplasms Working Group Consensus Recommendations for Diagnosis and Management of Primary Myelofibrosis, Polycytemia Vera, and Essential Thrombocytemia

Abstract
Myeloproliferative neoplasms (MPNs) are clonal disorders, derived from abnormal hematopoietic stem cells and result in an excessive production of blood cells. This MPN group of conditions encompasses different diseases with overlapping clinical and biologic similarities. The majority of the conventional therapies of MPN are palliative in nature. However, with the discovery of Janus Kinase 2 (JAK2) mutation and development of targeted JAK1/2 inhibition therapy, the therapeutic options in treatment landscape have changed dramatically. This article presents the revised Indian MPNs Working Group consensus recommendations. It highlights and brings into attention about the recent findings that have defined the state of the art of the diagnosis and therapy in the MPN area, including identification of the new driver and prognostic mutations, treatment goals in the management of myelofibrosis and polycytemia vera (PV), role of the recently approved, targeted tyrosine kinase inhibitor ruxolitinib in PV, and special issues such MPN consideration in patients with splenic vein thrombosis and the management of the disease in pregnancy.

Keywords: Essential thrombocytemia Janus kinase, myelofibrosis, myeloproliferative neoplasm, polycytemia vera

Introduction
Myeloproliferative neoplasms (MPNs) are a group of clonal disorders derived from multipotent hematopoietic myeloid progenitors.[1] The World Health Organization (WHO) classification of MPNs was updated in 2016 [Table 1] and has recognized polycytemia vera (PV), essential thrombocytemia (ET), and myelofibrosis (MF) as the classical Philadelphia chromosome (Ph)-negative or BCR-ABL1-negative MPNs.[2,3] Secondary MF can result from progression of PV and ET, with a 10-year risk of post-ET MF and post-PV MF being <4% and 10%, respectively.[4] All the 3 types of BCR-ABL1-negative MPNs pose a higher risk of transformation to acute myeloid leukemia (AML).[5] MPNs account for the large majority of myeloid malignancies[6] with incidence rates ranging from 0.1 to 3.1 cases/100,000/year in the United States and the European Union.[6-8] Among the 3 MPN subtypes, PV is the most common (0.4–2.8/100,000/year) compared to ET (0.38–1.7/100,000/year) and MF (0.1–1/100,000/year).[6-8] Epidemiological data on the incidence and prevalence of MPNs in the developing countries, including India, are sparse.

The cardinal features of the BCR-ABL1-negative MPN are the presence of an increased red cell mass in PV, an increase in the platelet counts in ET, and bone marrow fibrosis in MF.[2] The discovery of mutations such as Janus kinase 2 (JAK2) and thrombopoietin receptor (MPL) revolutionized the understanding of MPN pathophysiology and the development of targeted drug therapy in this area.[9]

A diverse array of somatic mutations, including driver mutations (JAK2 V617F, MPL, and calreticulin [CALR]) and high molecular risk prognostic mutations (EZH2, ASXL1, SRSF2, and IDH1), have been discovered recently, which led to a better understanding of the biology of MPNs and provided newer aspects of diagnosis, prognosis, and treatment.[10-12] Molecular biology assessment is an important component of the revised WHO diagnostic...
classification.[3] In addition to this, the treatment landscape of MPNs is also evolving with several targeted options being studied. As a result, in 2014, the MPN Working Group (MPN-WG) in India had worked on consolidating the consensus recommendations to improve diagnosis and prognostic stratification in patients with Ph-negative MPNs and published them, which have been well accepted by many of the practitioners in India.[13] With the recent updates in the literature, MPN-WG assembled to develop revised guidelines for the diagnosis and management of MPN based on the published literature and the experiences of the expert panel. MPN-WG experts’ panel will be called as panel throughout this manuscript.

Methods
The panel met with the objective of appraising the recent literature and revising the previously published consensus recommendations for the diagnosis and management of Ph-negative MPNs along with various other objectives focusing on raising awareness about MPNs through continuing medical education programs, conducting pathology workshops to improve diagnosis, and collaborating on MPN-related research. The methodology has been previously described.[13] The final version of the guidelines was further discussed in a consensus face-to-face meeting, which had included 2 international experts. The discussion also focused on some of the major challenges that were typically seen in Indian clinical practice. Appendix 1 describes how to appraise literature based on the level of evidence and hence, grades of recommendation.

Diagnosis of myeloproliferative neoplasm
The panel had agreed in the previous consensus meeting that WHO 2008 classification and diagnosis criteria for Ph-negative MPNs could be adapted in the Indian context with a few additional measures. Since there was a revision in the guidelines in 2016 the same has been updated in this revision statement [Table 1]. A detailed baseline workup has been presented in the previous version of the guidelines.[13] Based on these guidelines, diagnosis of MF, PV, and ET is currently a diagnostic algorithm that includes a composite assessment of biochemical and pathological assessments, and cytogenetic and mutational analysis. The laboratory detection of JAK2 V617F mutation, a highly sensitive and specific tool,[14] is the key pillar in the diagnosis of MPNs.

The diagnosis of MF is now divided into preprimary myelofibrosis (PMF) and overt MF. Overt MF diagnosis is primarily based on bone marrow fibrosis of Grade 2 or more, JAK2 V617F mutation positivity, progressive anemia, leukocytosis, splenomegaly, presence of constitutional symptoms, and leukoerythroblasticosis. Pre-PMF is diagnosed by above criteria with only 2 differences, bone marrow fibrosis grades below 2 and absence of leukoerythroblasticosis. In patients with JAK2 V617F-negative MF, MPL, and CALR mutation testing should be performed to further assist in diagnosis and to exclude other etiologies as causes of reactive thrombocytosis. The panel recommended that the BCR-ABL1 test should be performed to exclude chronic myelogenous leukemia. In the Indian settings, the panel recommended that the anemia related to MF should be confirmed by ruling out anemia due to any nutritional deficiencies or constitutional anemia. When marginal hemoglobin deficiency is observed, common nutritional causes should be diagnosed by simple blood tests and corrective measures, such as iron supplementation, which should be taken in patients where required.

The use of JAK2 V617F screening along with serum erythropoietin (EPO) level is particularly helpful in distinguishing PV from secondary erythrocytosis. Approximately, 3% of patients with PV who are negative for JAK2 V617F carry JAK2 exon 12 mutation.[15] The panel recommended incorporating JAK2 exon 12 mutation in the baseline workup of PV diagnosis. In line with the previous version of the consensus guideline, red cell mass measurement has not been recommended in diagnosing PV in the routine practice. As discussed in the previous version of the guidelines, there is a paucity of specialized bone marrow biopsy centers in India; therefore, distinguishing ET from prefibrotic PMF-based on the bone marrow morphology criteria would be challenging in India. In this case, minor criteria of the WHO guidelines can be used to confirm histologic pattern indicative of prefibrotic PMF.[16] The panel suggested that pathology slides should be referred to MPN reference centers to establish and confirmed the diagnosis.
### Table 1: Diagnostic criteria for myeloproliferative neoplasm (adapted from World Health Organization 2016 diagnostic criteria)

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Description</th>
</tr>
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<tbody>
<tr>
<td><strong>WHO pre-PMF criteria</strong></td>
<td><strong>Major criteria</strong></td>
</tr>
<tr>
<td></td>
<td>Megakaryocytic proliferation and atypia, without reticulin fibrosis greater than Grade 1, accompanied by increased age-adjusted BM cellularity, granulocytic proliferation, and often decreased erythropoiesis Not meeting the WHO criteria for BCR-ABL1 positive CML, PV, ET, MDSs, or other myeloid neoplasms Presence of JAK2, CALR, or MPL mutation or in the absence of these mutations, presence of another clonal marker (ASXL1, EZH2, TET2, IDH1/IDH2, SRSF2, SF3B1), or absence of minor reactive BM reticulin fibrosis</td>
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<td></td>
<td><strong>Minor criteria</strong></td>
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<tr>
<td></td>
<td>Presence of at least 1 of the following, confirmed in 2 consecutive determinations</td>
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<tr>
<td></td>
<td>Anemia not attributed to a comorbid condition</td>
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<td></td>
<td>Leukocytosis ≥11x10⁹/L</td>
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<tr>
<td></td>
<td>Palpable splenomegaly</td>
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<tr>
<td></td>
<td>LDH increased to above upper normal limit of institutional reference range</td>
</tr>
<tr>
<td><strong>Diagnosis</strong></td>
<td>Diagnosis of pre-PMF requires meeting all 3 major criteria, and at least 1 minor criterion</td>
</tr>
<tr>
<td><strong>WHO overt PMF criteria</strong></td>
<td><strong>Major criteria</strong></td>
</tr>
<tr>
<td></td>
<td>Presence of megakaryocytic proliferation and atypia, accompanied by either reticulin and/or collagen fibrosis Grades 2 or 3</td>
</tr>
<tr>
<td></td>
<td>Not meeting WHO criteria for ET, PV, BCR-ABL1 CML, MDSs, or other myeloid neoplasms</td>
</tr>
<tr>
<td></td>
<td>Presence of JAK2, CALR, or MPL mutation or in the absence of these mutations, presence of another clonal marker, or absence of reactive myelofibrosis</td>
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<tr>
<td></td>
<td><strong>Minor criteria</strong></td>
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<tr>
<td></td>
<td>Presence of at least 1 of the following, confirmed in 2 consecutive determinations</td>
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<tr>
<td></td>
<td>Anemia not attributed to a comorbid condition</td>
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<tr>
<td></td>
<td>LDH increased to above upper normal limit of institutional reference range</td>
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<tr>
<td></td>
<td>Leukoerythroblastosis</td>
</tr>
<tr>
<td><strong>Diagnosis</strong></td>
<td>Diagnosis of overt PMF requires meeting all 3 major criteria, and at least 1 minor criterion</td>
</tr>
<tr>
<td><strong>WHO PV criteria</strong></td>
<td><strong>Major criteria</strong></td>
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<tr>
<td></td>
<td>Hb &gt;16.5 g/dL in men &gt;16.0 g/dL in women or Hematocrit &gt;49% in men &gt;48% in women or Increased RCM BM biopsy showing hypercellularity for age with trilineage growth (panmyelosis) including prominent erythroid, granulocytic, and megakaryocytic proliferation with pleomorphic, mature megakaryocytes (differences in size) Presence of JAK2V617F or JAK2 exon 12 mutation</td>
</tr>
<tr>
<td></td>
<td><strong>Minor criteria</strong></td>
</tr>
<tr>
<td></td>
<td>Subnormal serum erythropoietin level</td>
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<tr>
<td><strong>Diagnosis</strong></td>
<td>Diagnosis of PV requires meeting either all 3 major criteria, or the first 2 major criteria and the minor criterion'</td>
</tr>
<tr>
<td><strong>WHO ET criteria</strong></td>
<td><strong>Major criteria</strong></td>
</tr>
<tr>
<td></td>
<td>Platelet count ≥450x10⁵/L BM biopsy showing proliferation mainly of the megakaryocyte lineage with increased numbers of enlarged, mature megakaryocytes with hyperlobulated nuclei. No significant increase or left shift in neutrophil granulopoiesis or erythropoiesis and very rarely minor (Grade 1) increase in reticulin fibers Not meeting WHO criteria for BCR-ABL1 CML, PV, PMF, MDSs, or other myeloid neoplasms Presence of JAK2, CALR, or MPL mutation</td>
</tr>
<tr>
<td></td>
<td><strong>Minor criteria</strong></td>
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<tr>
<td></td>
<td>Presence of a clonal marker or absence of evidence for reactive thrombocytosis</td>
</tr>
<tr>
<td><strong>Diagnosis</strong></td>
<td>Diagnosis of ET requires meeting all 4 major criteria or the first 3 major criteria and the minor criterion</td>
</tr>
</tbody>
</table>

PMF – Primary myelofibrosis; WHO – World Health Organization; BM – Bone marrow; CML – Chronic myeloid leukemia; PV – Polycythemia vera; ET – Essential thrombocythemia; Hb – Hemoglobin; MDSs – Myelodysplastic syndromes; JAK2 – Janus kinase 2; CALR – Calreticulin; LDH – Lactate dehydrogenase; RCM – Red cell mass
Recently, a new algorithm “reflex test” has been proposed. This reflex test sequentially evaluates for the common major gene mutations associated with MPNs. In this test, the presence of JAK2 V617F mutation is determined as a first step. If this result is negative, assessment for CALR mutations is performed. If the CALR result is also negative, then testing proceeds to evaluate for mutations in exon 10 of the MPL gene. If either JAK2 V617F or CALR mutations are detected in the process, the testing algorithm ends. Panel discussed the following typical order of testing of genetic mutations: JAK2, CALR, cMPL, and JAK2 exon 12 (in patients with MF); JAK2, JAK2 exon 12, cMPL, and CALR (in patients with PV); and JAK2, cMPL, CALR, and JAK2 exon 12 (in patients with ET).[17]

**Myelofibrosis**

The clinical manifestations of MF are heterogeneous and up to 30% of the patients are initially asymptomatic, and as the disease evolves, patients become symptomatic with an varied spectrum of disease presentation (bone marrow fibrosis, splenomegaly causing abdominal symptoms and early satiety, and constitutional symptoms such as weight loss, night sweats, and low-grade fever, aquagenic pruritus, bone pain, or thrombosis). With the exception of allogeneic stem cell transplantation, currently available drug therapies for MF are not curative, and the treatment choice is mainly guided by the symptom or feature or risk stratification [Figure 1].[18] The conventional treatment for MF are directed to treat anemia and other cytopenias, to manage symptomatic splenomegaly and constitutional symptoms (weight loss, night sweats, fever, pruritus, fatigue, etc.), to improve quality of life, and to improve survival and minimize the risk of transformation to acute leukemia. Other possible goal is to avoid first occurrence or recurrence of thrombotic and bleeding complications.[19,20]

**Risk stratification and assessing prognosis**

The International Prognostic Scoring System (IPSS) is the most commonly used prognostic model for determining the survival risk category at the time of diagnosis. Based on the IPSS, patients with PMF are diagnosed as having low-, intermediate 1-, intermediate 2-, or high-risk disease. IPSS uses the following 5 risk factors to predict prognosis and assign a patient to a risk group: (1) age >65 years; (2) hemoglobin <10 g/dL; (3) leukocyte count >2.5 × 10⁹/L; (4) circulating blood blasts >1%; and (5) the presence of constitutional symptoms.[21] In addition, dynamic prognostic models (dynamic IPSS [DIPSS] and DIPSS Plus) are available based on the acquisition of additional risk factors during the disease course that may substantially modify patients’ outcomes.[22,23] The DIPSS uses the same 5 risk factors and has been validated to predict prognosis at any time during the disease course. The DIPSS was recently modified (DIPSS Plus) with the incorporation of the following 3 additional risk factors: (1) red blood cell transfusion need; (2) platelet count <100 × 10⁹/L; and (3) unfavorable karyotype.[22]

**Stem cell transplantation**

Multiple studies have confirmed that allogeneic stem cell transplant is the only curative treatment in PMF. As this is a disease of elderly patients, only a small minority of patients are candidates for SCT. It is generally considered only for patients with intermediate-2 or high-risk disease (patients with an expected survival of <5 years) due to the high procedure-related morbidity and mortality.[21-23] A transplant-eligible patient is defined as one deemed fit enough to undergo the procedure with manageable comorbidities and having an HLA-matched sibling or unrelated donor available. Patients above 45 years have a very poor survival on myeloablative conditioning.[24]

**Recommendation**

There are no randomized controlled trials comparing allo-HSCT to any alternative/supportive therapy. Allogeneic stem cell transplant with myeloablative or reduced intensity conditioning is indicated in young (<40 years of age), Int-2, or high-risk patients with PMF. Reduced-intensity transplantation should be considered for patients aged 40–60 years with Int-2 or high risk at diagnosis or later during the disease. Grade B recommendation, evidence Level III.[19]

**Management of anemia**

**Erythropoietin**

The use of recombinant human erythropoetin (rEPO) appears to be limited principally to a subgroup of patients with inappropriately low endogenous EPO levels (<200 IU) and with relatively moderate anemia.[19,25] EPO levels should be measured before starting therapy with recombinant human EPO. The targeted hemoglobin level
is 10 g/dL in patients with MF. EPO should be initiated at a dose of 10,000 U once weekly, and response should be monitored for 2 weeks, and if no response is observed, the dose can be escalated up to 40,000 IU once weekly. Treatment should be discontinued after 3–4 months if no response occurs. Darbepoetin has greater activity and a longer half-life than EPO, which allows for less frequent administration. Darbepoetin-α, administered bimonthly, is equally effective, and the recommended dose is in the range of 150–300 µg/fortnight. Recombinant human EPO should be stopped or dose reduced when hemoglobin is above 12 g/dL to avoid the risk of thrombosis.[19] In a recent analysis, the combination of EPO with ruxolitinib was evaluated. In this study, clinical benefits were observed in patients treated with ruxolitinib. Moreover, recently a sub-analysis of pooled data of COMFORT trials has shown that the postbaseline anemia induced by ruxolitinib in the treatment of MF has no prognostic impact on the overall survival (OS) of the patients, and the anemia due to ruxolitinib is manageable with conventional treatments than the disease-related baseline anemia, which is an established poor prognostic factor.[26]

Recommendation

EPO is recommended as the first-line therapy for the treatment of anemia in PMF only for patients with EPO levels <200 IU.[19,20] Grade B recommendation, evidence Level III.[24]

Danazol

Danazol, a synthetic attenuated androgen, has been considered as the first-line androgen of choice in MF. Recommended starting dose is 200 mg daily, with a gradual dose escalation, depending on tolerability and patient weight (to a maximum of 600 mg daily for patients <80 kg and 800 mg for patients >80 kg). Most patients respond within the first 2–3 months. The therapy should be discontinued if no response is observed after a 4-month trial of these agents. While on danazol treatment, liver function of patients should be monitored regularly (once monthly), and the male patients should be screened for prostate cancer. A synergistic effect between human recombinant EPO and danazol treatment has been reported.[19,27]

Recommendation

Danazol is recommended as an alternative first-line therapy in the treatment of anemia in patients with PMF associated with transfusion-dependent anemia.[19,27] Grade B recommendation, evidence Level III.[19,27]

Thalidomide

Low-dose thalidomide (50 mg/day preferably given at bedtime) can be used for the treatment of patients with MF with anemia and thrombocytopenia, and for the reduction of splenomegaly.[28] Low-dose thalidomide in combination with prednisolone showed significant responses for anemia (67%), thrombocytopenia (75%), and splenomegaly (33%).[29] However, thalidomide is associated with nonhematological toxicity (constipation, sedation, depression, and peripheral neuropathy) and is contraindicated in pregnancy. It is to be used with caution and with effective counseling and contraception in women of childbearing age.

Recommendation

Low-dose thalidomide (maximum 50 mg/day) in combination with prednisolone (10–20 mg/day for 2 weeks and afterward tapering to the lowest dose necessary for maintaining an adequate hemoglobin concentration) can be considered for patients not responding to EPO or danazol. Therapy should be discontinued if no response is seen after 4 months. Grade B recommendation, evidence Level III.[19]

Glucocorticoids (mostly used in combination with thalidomide)

Corticosteroids, especially prednisone, have been used for the palliation of constitutional symptoms and anemia in patients with MF. They are recommended chiefly in patients with hemolytic activity, but could be considered at a dose of 10–20 mg/day in some patients with anemia without evidence of hemolysis. In patients with hemolysis and a positive Coombs test, an initial test dose of 1 mg/kg is recommended. Grade C recommendation, evidence Level IV.[19]

The panel agreed that standard transfusion guidelines should be applied for patients eligible for transfusion to correct anemia associated with MF.

Treatment of thrombocytopenia

The panel has examined the available evidence for the treatment of thrombocytopenia associated with MF and concluded that in the context of hematology practice in India, thalidomide in combination with prednisolone should be the choice of therapy.[17] If no response is observed with this combination, danazol is recommended.

Treatment of symptomatic splenomegaly

Hydroxyurea

Hydroxyurea is one of the most commonly used agents despite the fact that the available data are limited to support its efficacy in MPN. Based on the prior experience of the panel, the members were in agreement that they have seen only modest improvement in symptoms with hydroxyurea.[30,31]

Recommendation

Hydroxyurea (0.5–2 g/d) is recommended as cytoreductive therapy (CYTO) in older patients with MF who are not eligible for transplantation. Grade B recommendation, evidence Level III.[19]

Janus kinase 2 inhibitors

JAK inhibitors are the only drugs that have been evaluated in the randomized clinical trials. A number of JAK
inhibitors are at various stages of clinical development, and a consistent pattern of response in splenomegaly and disease-related symptoms is emerging.

In India, ruxolitinib is approved for the treatment of patients with MF, including primary MF, post-PV MF or post-ET MF. Ruxolitinib monotherapy has been evaluated in intermediate-2- or high-risk IPSS patients with MF in phase-III COMFORT studies which have completed 5 years long-term follow-up. In COMFORT-II, 97.1% of patients who remained on treatment had improvements from baseline in spleen volume, and 53.4% achieved ≥35% reductions. Spleen reductions of ≥35% among ruxolitinib randomized patients were sustained with long-term therapy (median duration, 3.2 years); the probability of maintaining a spleen response was 0.48 (95% confidence interval [CI], 0.35–0.60) at 5.0 years for patients initially randomized to ruxolitinib. In addition, long-term findings from COMFORT studies support the disease-modifying effect of ruxolitinib. Among patients randomized to ruxolitinib in COMFORT-II, 23 (15.8%) had improved fibrosis (any grade), 47 (32.2%) had stable fibrosis at their last assessment (median treatment duration, 2.2 years in the ruxolitinib arm). Patients randomized to ruxolitinib had longer OS compared than those randomized to best available therapy (BAT), with a 33% reduction in risk of death with ruxolitinib treatment (HR, 0.67 [95% CI, 0.44–1.02]; P = 0.06). Similar results were also found in COMFORT-I, for all ruxolitinib treated pts (n = 57), 33% (G[-1], n = 11; G [-2], n = 7; G [-3], n = 1) had an improvement, 49% had no change or stabilization, and 18% had a worsening in BM fibrosis from BL to the last grade. A pooled OS analysis of 5-year data from the COMFORT-I and COMFORT-II shows that the risk of death was reduced by 30% among patients randomized to ruxolitinib compared with control patients (median OS: ruxolitinib, 5.3 years; control, 3.8 years; hazard ratio [HR; ruxolitinib vs. control], 0.70; 95% CI, 0.54 – 0.91; P = 0.0065;) as per intent-to-treat analysis and it was more pronounced after censoring patients at the time of crossover, demonstrated that ruxolitinib prolonged survival compared with control (median OS: ruxolitinib, 5.3 years; control, 2.4 years; HR [ruxolitinib vs. censored at crossover], 0.53; 95% CI, 0.36 – 0.78; P = 0.0013). The reasons for the survival advantage observed with ruxolitinib may be due to the number of benefits of treatment in terms of spleen size reduction, alleviation of cytokine-driven symptoms and inflammation, improvement of overall nutritional status, and reduced fibrosis in some patients. Ruxolitinib is the current treatment of choice for constitutional prognostic symptoms of the disease (weight loss, fever, and night sweats), spleen-related symptoms (such as abdominal discomfort and pain under left ribs), and disease-related symptoms (such as itching, fatigue), as no other therapy has been shown to significantly improve symptom burden and quality of life in MF. Based on these benefits including reduction of symptom burden, NCCN guidelines in the management of MPNs has recommended the use of ruxolitinib in low risk/int-1/int-2 and high-risk patients with symptoms. A common misconception is that JAK inhibition is effective primarily in patients who have the JAK2 V617F mutation. However, ruxolitinib demonstrated the comparable efficacy in patients with or without the V617F mutation. Since JAK2 is involved in thrombopoietin and EPO signaling, dose-dependent thrombocytopenia and anemia should be anticipated and managed appropriately in patients with MF undergoing treatment with JAK2 inhibitors such as ruxolitinib. Recent updates from COMFORT studies have shown that the postbaseline anemia induced by ruxolitinib in the treatment of MF does not have an impact on the OS of the patients, and it can be managed with conventional treatments than the disease-related baseline anemia, which is an established poor prognostic factor. Starting dose of ruxolitinib should be personalized according to baseline platelet counts [Table 2] and there is no separate guidance for elderly patients since both groups showed comparable clinical responses and toxicity rates similar to younger individuals, however, comorbidities have to be kept in mind while treating elderly patients with MF.

Table 2: Ruxolitinib dose in primary myelofibrosis

<table>
<thead>
<tr>
<th>Platelet count</th>
<th>Starting dose</th>
</tr>
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<tbody>
<tr>
<td>&gt;200×10^9/L</td>
<td>20 mg orally bid</td>
</tr>
<tr>
<td>100-200×10^9/L</td>
<td>15 mg orally bid</td>
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<tr>
<td>50–&lt;100×10^9/L</td>
<td>5 mg orally bid</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Dose modifications (with starting platelet count of at least 100×10^9/L)</th>
<th>Maximum dose based on platelet count after prior treatment interruption or dose reduction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current platelet count</td>
<td>Dose at time of decline in platelet count (new dose to be used)</td>
</tr>
<tr>
<td>20 mg bid</td>
<td>15 mg bid</td>
</tr>
<tr>
<td>≥125×10^9/L</td>
<td>No change</td>
</tr>
<tr>
<td>100–&lt;125×10^9/L</td>
<td>15 mg bid</td>
</tr>
<tr>
<td>75–&lt;100×10^9/L</td>
<td>10 mg bid</td>
</tr>
<tr>
<td>50–&lt;75×10^9/L</td>
<td>5 mg bid</td>
</tr>
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</table>

Bid – Twice daily

Recommendation

Ruxolitinib should be considered as the first choice of treatment in patients with symptomatic and progressive...
splenomegaly. Grade A recommendation, evidence Level Ib.[19,39]

**Splenectomy**

Even though splenectomy procedure is associated with significant morbidity (25%-30%) and mortality (7%-10%), in a restricted subgroup of patients with MF associated with refractory splenomegaly, this option should be carefully evaluated. Significant surgical expertise, fastidious surgical hemostasis, and careful control of postoperative thrombocytosis have been found to be essential in trying to minimize the risk of this procedure. Critical factors to be assessed before splenectomy include the type of conditioning used, timing of the intervention, and expertise of the surgeon.[27]

**Recommendation**

Splenectomy should not be considered for symptomatic splenomegaly. Splenectomy should be considered only in patients who are not responsive to hydroxyurea, interferon-α (IFN-α), and/or ruxolitinib, with marked splenomegaly associated complications such as splenic infarct, splenic abscess, or rupture, repeated upper gastrointestinal bleeding episodes due to portal hypertension and/or cytopenias secondary to hypersplenism. Grade B recommendation, evidence Level III.[39]

**Splenic irradiation**

Splenic irradiation should be reserved for patients in the indications mentioned above who are not responsive to conventional and novel therapies and patients who are not eligible for splenectomy. Grade B recommendation, evidence Level III.[19] Splenic embolization is not recommended for symptomatic splenomegaly in MF.

**Treatment of symptoms**

Ruxolitinib is the current treatment of choice, as no other therapy has been shown to significantly improve symptom burden and quality of life in MF. Grade A recommendation, evidence Level Ib.[19]

**Combination therapy**

As further details of molecular mechanisms leading to Ph-negative MPNs emerge, combinations of targeted agents are under investigation to improve outcomes in these patients. In preclinical studies, pan histone deacetylase inhibitor, panobinostat has shown synergistic effect when combined with ruxolitinib, especially due to deacetylation of protein chaperone HSP90 involved in JAK-STAT signaling.[28] Preliminary results from a phase1b study in patients with MF have provided supportive evidence for the combination treatment strategies in MF.[29] However, such therapy should only be considered in the clinical trial setting.

**Avoiding thrombotic and bleeding complications**

Retrospective analyses indicate that the incidence of thrombotic complications were similar in PMF and ET.[32,33] No prospective trials of platelet-reducing agents or aspirin have been performed in PMF. The panel suggested that clinicians should follow the guidelines provided for ET regarding thrombosis and bleeding prevention in PMF. Other factors contributing to atherosclerosis should be addressed. General care guidelines for preventing thrombosis should be followed.

**Evaluation of response and follow-up**

Clinical assessments are recommended by the panel at regular intervals to monitor response to therapy. The panel has updated their recommendation for the use of revised International WG-MPNs Research and Treatment and the European Leukemia Net (ELN) criteria. The definition of response as per the criteria should be restricted to clinical trial settings only.[40]

**Polycythemia vera**

**Treatment goals**

The goals of current therapies in PV are to prevent the occurrence of thrombosis/vascular events; delay transformation to MF or AML; reducing constitutional symptoms; maintaining hematocrit <0.45, and managing special situations such as pregnancy or cardiovascular risk. In a large-scale, prospective, randomized clinical trial (CYTO in PV study [n = 365]), the benefit of maintaining hematocrit <45% versus 45%-50% using conventional treatment was associated with a lower rate of cardiovascular deaths and major thrombotic events in patients with a target hematocrit of <45% than in those with a target hematocrit of 45%-50%. Revised WHO criteria-2016 have been able to better identify patients with masked PV (mPV). Patients with mPV, who tend to have normal or borderline hematocrits values but are usually positive for JAK2 mutations, have bone marrow features consistent with PV, and have low serum EPO levels. The revision of the WHO diagnostic criteria with emphasis on a lower hemoglobin threshold and/or the use of hematocrit threshold values may be helpful in accurately diagnosing those with mPV and could allow for appropriate and prompt treatment of these patients.[3,42]

**Risk stratification**

Therapeutic options in PV are limited, and no cure is available. Initial treatment depends on the risk stratification of the patient. Patients can be stratified in high- or low-risk categories according to whether they are older or younger than 60 years and have a history of thrombosis. Low-risk patients have zero risk factors; high-risk patients have 1 or 2 risk factors. Leukocytosis and JAK2 V617F allele burden have been identified as other key risk factors.[43]

**Clinical management**

The panel examined the clinical evidence for traditional treatments approaches in PV, and the recommendations
were formulated based on the consensus. The treatment algorithm for PV is provided in Figure 2.

**Phlebotomy**

The objective of phlebotomy in PV is to improve the circulation of blood by lowering the blood viscosity. The basis of conventional therapy is to maintain hematocrit at <0.45 and <0.42 in male and female patients, respectively. This goal can be achieved in low-risk patients with phlebotomy. Phlebotomy should be performed by removing 250–500 mL of blood every other day or twice a week, until the target hematocrit is achieved. Patients with impaired cardiovascular function are better treated with smaller phlebotomies at more frequent intervals. Phlebotomy is recommended for hematocrit levels >0.48. Grade A recommendation, evidence Level Ib.[19]

**Aspirin therapy**

For all patients with PV, low-dose aspirin (i.e. 100 mg daily or 81 mg in the US) is recommended. In a large randomized study, aspirin has shown superiority over placebo with a significant reduction in cardiovascular death, thrombosis, strokes, and in overall mortality.[45] Aspirin has been shown to reduce both arterial and venous thrombosis in PV.[41] Aspirin should not be given to patients with platelets more than 1500 × 10⁹/L due to an increased risk of bleeding, instead CYTO should be initiated. In case of aspirin allergy, clopidogrel can be used. Grade A recommendation, evidence Level Ib.[19]

**Choice of cytoreductive therapy in polycythemia vera**

**Interferon-α**

Although the supporting evidence for IFN-α efficacy in PV is limited, hydroxyurea is the recommended first-line treatment.

*Figure 2: Algorithm for management of essential thrombocytopenia*
treatment for high-risk patients. Pegylated IFN-α 2a (PEG IFN-α 2a) has shown better clinical response rates and demonstrated more activity with few complete molecular remissions (as determined by JAK2 allele burden) at a better side effect profile than non-PEG versions of IFNα.[44]

**Recommendation**

IFN-α can be considered in younger patients (<40 years). For patients >40 years of age, hydroxyurea should be considered as the first-line of treatment. The panel recommends that the lower doses of IFN should be used in India because of smaller body mass index as compared to Western population and hence, the dose of PEG IFN in Indian patients may range from 40, 80, to 150 µg, weekly. Grade B recommendation, evidence Level III.[19]

**Hydroxyurea**

Hydroxyurea is recommended as a first-line CYTO in patients with PV >40 years or younger patients who do not tolerate IFN-α (Grade A recommendation, evidence Level Ib).[43] Hydroxyurea is useful in controlling PV-related symptoms; splenomegaly, leukocytosis, thrombocytosis, and hematocrit. However, hydroxyurea-treated patients can eventually become resistant or experience unacceptable adverse effects (hydroxyurea intolerance), including skin ulcers, a reduction in blood cells, gastrointestinal problems, oral ulcers, stomatitis, hyperkeratosis, or actinic keratosis.[46] However, hydroxyurea should be continued until patients develop intolerance or progression.[43] Increased risk of leukemic transformation has not been observed despite long-term hydroxyurea treatment. Grade A recommendation, evidence Level Ib. The panel recommended maximum tolerated dose of hydroxyurea as 1 g (this is with reference to 2 g in hydroxyurea resistance/tolerance ELN criteria).

**Busulfan**

Busulfan can be used as the second-line CYTO for elderly population of age >70 years. Sequential use of busulfan and hydroxyurea resulted in a significant increase in the risk of second malignancies, including leukemias.[47]

**Janus kinase 2 inhibitors**

Like MF landscape, several JAK inhibitors are under evaluation in clinical trials for efficacy and safety in patients with PV. Ruxolitinib is the only JAK1/2 inhibitor approved by the United States Food and administration and European Medicine Agency for the treatment of patients with PV who have had an inadequate response to or are intolerant of hydroxyurea. Findings from the pivotal RESPONSE study showed that ruxolitinib provided clinically relevant improvements compared with BAT in maintaining control of hematocrit level without phlebotomy, normalizing blood cell count, reducing spleen volume, and improving symptoms. In PV patients with inadequately controlled leukocyte count, recent evidence has shown that ruxolitinib treatment resulted in better white blood cell (WBC) control as compared to conventional therapies such as hydroxyurea (HU) and BAT.[48] Recent data from RESPONSE-II, a trial conducted in PV patients without splenomegaly, have also shown hematocrit control, complete hematologic response, and symptom control among ruxolitinib-treated patients better than BAT.[49] In addition, there were fewer thromboembolic events in the ruxolitinib arm as compared to BAT during the therapy. There is also some preliminary data on the reduction of JAK2 allele burden by ruxolitinib up to −22% by week 80; however, the impact of this on the disease has to be studied further. The rates of all Grade and Grade 3/4 thromboembolic events per 100 patient-years of exposure were lower among patients originally randomized to ruxolitinib than BAT. The starting dose of ruxolitinib in RESPONSE was 10 mg twice daily.[48]

**Recommendation**

Ruxolitinib should be considered as first-choice of treatment in patients with PV who have had an inadequate response to or are intolerant of hydroxyurea. Proposed definition of resistance/intolerance to HU was defined as a dose ≥2 g/day or a maximum tolerated dose <2 g/day resulting in at least 1 of the following: Need for phlebotomy to maintain HCT <45%; Platelet count >400 × 10⁹/L and WBC count >10 × 10⁹/L; Failure to reduce splenomegaly extending >10 cm below the costal margin by >50%, as measured by palpation; Absolute neutrophil count <1.0 × 10⁹/L or platelet count <100 × 10⁹/L or hemoglobin <100 g/L at the lowest dose of HU required to achieve a complete or partial response; Presence of leg ulcer or unacceptable HU-related nonhematologic toxicities.[46] Grade A recommendation, evidence Level Ib.[19]

**Anagrelide**

Anagrelide has powerful platelet reducing activity that could be helpful in the management of patients who are intolerant or refractory to hydroxyurea, INF-α, or ruxolitinib. Grade C recommendation, evidence Level IV.[19]

**Radioactive phosphorus**

Due to its leukemogenic effect, radioactive phosphorus is not a recommended option for patients <75 years.[19]

**Evaluation of response and follow-up**

The panel recommends that the goal of therapy should be normalization of peripheral blood counts, and the ELN criteria for PV should be used for the evaluation of response in clinical trial setting.

**Essential thrombocythemia**

**Treatment goals**

Similar to PV, the main treatment goals in ET include minimizing the risk for thrombosis and progression, normalizing peripheral blood counts, reducing constitutional
symptoms, and managing special situations like pregnancy. Treatment algorithm recommended for Indian scenario is discussed in the Figure 3.

**Risk stratification**

Risk stratification in ET is based on the assessment of risk of thrombosis, as the current therapy in ET is aimed at lowering the risk of thrombosis. True ET diagnosed, according to the 2008 WHO classification, has not been reported to affect the life expectancy of patients. High-risk is defined as the presence of age above 60 years or history of previous thrombosis, or a platelet count more than $1500 \times 10^9/L$; and these patients should be treated with CYTO. Low-risk is defined by the absence of these factors and should not be treated with CYTO except in patients with uncontrolled cardiovascular risk factors.

**Clinical management**

Patients with ET are at high risk for thrombosis; hence, vigorous treatment is required for managing cardiovascular risk factors. Hence, cardiovascular risk assessment is required and will dictate the overall management of the patient. It is important to emphasize that before starting therapy, patients should be evaluated for eventual progression to MF if they show symptomatic or progressive splenomegaly, other evidence of disease progression such as weight loss, night sweats, progressive leukocytosis, and/or thrombocytosis. Stem cell transplantation is almost never performed in ET due to unfavorable risk-benefit profile. The treatment algorithm for ET is provided in Figure 3.

**Aspirin**

Antiplatelet therapy with aspirin 75–150 mg/day is recommended, unless otherwise contraindicated for all the patients with ET. Aspirin should not be given to patients with platelets more than $1500 \times 10^9/L$ due to an increased risk of bleeding; instead CYTO should be initiated; recommendation, evidence Level IIIB.

**Choice of cytoreductive therapy in essential thrombocythemia**

**Hydroxyurea**

Hydroxyurea is the best-documented therapy in ET and is recommended as a first-line therapy in the majority of patients with high-risk ET. Hydroxyurea markedly reduces thrombotic complications compared to aspirin alone. Panel recommended that patients presenting a symptom score of <8 points should be closely followed up every 3 months for the progression of symptoms.

**Interferon-α**

IFN-α treatment is well documented and safe in ET and is not considered leukemogenic or teratogenic. PEG IFN has also been used; Level IB evidence to support the use of IFN. IFN-α is the recommended first-line therapy in younger patients. It can be used in older patients if long-term use of hydroxyurea is not suitable and in patients who do not tolerate hydroxyurea (Grade B recommendation, evidence Level III). IFN-α is the treatment of choice if CYTO is indicated during pregnancy or when pregnancy is planned.

**Anagrelide**

This may be used as the second-line treatment in patients resistant to or intolerant of hydroxyurea. It has been found to be equivalent to hydroxyurea although some studies have reported higher risk of thrombosis.

**Janus kinase 2 inhibitors**

Since no JAK2 inhibitor has been studied extensively in ET to date, these drugs are experimental at the present time and should not be used outside of the context of clinical trials.

**Other drugs**

Other drugs such as busulfan and radioactive phosphorus are not relevant in ET. Anagrelide has many complications, but may be used in some patients, if necessary, in combination with hydroxyurea to control platelet counts.

**Management of complications of myeloproliferative neoplasms**

**Acute thrombotic events and secondary prophylaxis**

In general, acute thrombotic events should be treated as in non-MPN patients. Control of hematocrit and platelet count
should be optimized. In emergency situations such as acute cerebrovascular complications or severe digital ischemia, acute platelet apheresis or erythropheresis can be used in order to achieve a rapid reduction in blood counts. Since the effect is brief, CYTO preferably with hydroxyurea should be started as soon as possible in patients not on CYTO. Prevention of recurrent thrombosis should be independently achieved in patients with previous venous thrombosis by both oral anticoagulants and antiplatelet drugs. Since no prospective trials exist, it remains unclear whether it is better to give a short course of warfarin or to continue with long-term therapy for secondary prevention of venous thromboembolism.

**Bleeding**

The most important cause of bleeding in ET and PV is acquired von Willebrand syndrome associated with high platelet counts (>1500 × 10^9/L). Therefore, the most important therapeutic intervention to manage acute bleeding in the patients with thrombocytopenia is platelet reduction, and the recommended agent is hydroxyurea.

Platelet apheresis is indicated when extreme thrombocytosis is accompanied by an urgent need to reduce platelet counts, i.e., severe or life-threatening bleeding.

**Transformation to acute myeloid leukemia**

The results after conventional induction chemotherapy are dismal in patients developing AML after PV, ET, or PMF, with a very short median survival. If possible, it is recommended that patients undergo allo-SCT after induction chemotherapy. Hypomethylating agents can be used as bridge to transplant.

**Splanchnic vein thrombosis and portal vein hypertension**

Splanchnic vein thrombosis (SVT) includes the Budd-Chiari syndrome (BCS) and portal vein thrombosis (PVT). MPNs are the most frequent underlying prothrombotic factors in BCS and PVT. Peripheral blood cell counts often remain within a normal range because of portal hypertension and its sequelae (splenomegaly, hemodilution, and iron deficiency). In a meta-analysis, high prevalence of JAK2 V617F in SVT was reported. The strong relation between MPNs and SVT is confirmed by the high prevalence of JAK2V617F in these patients. It is suggested that JAK2 V617F screening should be included in the routine diagnostic workup of all patients with SVT, regardless of the absence of MPN hallmarks, such as elevated peripheral blood counts.

**Special issues**

**Pregnancy**

There is limited information in the medical literature about the management of MPNs in pregnancy. The live birth rate is about 60% due to an overall incidence of first trimester miscarriage of 31%–36% (about twice the normal rate) and an increased risk of intrauterine growth retardation, intrauterine death, and stillbirth (8%). Major maternal complications are less common and occur in approximately 8% of patients with ET.

Pregnancy is likely to be accompanied by a high risk of complications for the mother and/or fetus if any of the following factors are present: previous venous or arterial thrombosis in mother, previous hemorrhage attributed to PV/ET, previous pregnancy complications that may have been caused by PV/ET (these include significant antepartum hemorrhage or postpartum hemorrhage, severe preeclampsia, unexplained recurrent first trimester loss (≥3), intrauterine growth retardation (<5 percentile for gestation), intrauterine death or stillbirth with no other cause identified, placental abruption), and platelet count above 1000 × 10^9/L.

Therapeutic options include antithrombotic treatment, phlebotomy in PV, and cytoreductive agents. The optimal hematocrit in pregnancy has yet to be established, but the current recommendation is to maintain the hematocrit within the normal range appropriate for gestation. The increased plasma volume often results in a reduced hematocrit and platelet count during the second trimester. The levels rise again during the postpartum period contributing to an increased thrombotic risk during the first 6 weeks after delivery. Close monitoring of blood counts is important during this period. Low-dose aspirin is safe and seems advantageous during pregnancy in ET. Starting on the day of the delivery, aspirin is substituted by a prophylactic dose of low molecular weight heparin (LMWH), which is given until 6 weeks after delivery. The doses of LMWH that have been reported are dalteparin 5000 U or enoxaparin 40 mg daily.

**Pediatric myeloproliferative neoplasm**

The incidence of different MPN in patients aged <20 years is so low that formal evidence-based recommendations are impossible to provide.

**Discussion**

The diagnostic criteria and treatment landscape of MPNs are evolving at a fast pace. In this paper, the panel has appraised the recent literature and updated the existing consolidated MPN guidelines to further strengthen the understanding of fellow practitioners. The recommendations are developed solely in consensus with the experts in the field of Ph-negative classical MPNs, and a group decisionmaking process was adopted while interpreting available clinical evidence in the context of hematology practice in India. Improved diagnostic capabilities among the Indian practitioners will in turn help in identifying the true incidence and prevalence of the disease. In addition, improved diagnosis with rational treatment will help in achieving better treatment outcomes in these patients, who were previously left undiagnosed because of the insidious nature of the disease.
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Conflicts of interest
There are no conflicts of interest.

References