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Recent advances in the genomics and therapy of *BCR/ABL1*-positive and -negative chronic myeloproliferative neoplasms

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ABSTRACT

This review is based on the presentations and deliberations at the 7th John Goldman Chronic Myeloid Leukemia (CML) and Myeloproliferative Neoplasms (MPN) Colloquium which took place in Estoril, Portugal on the 15th October 2017, and the 11th post-ASH International Workshop on CML and MPN which took place on the 6th-7th December 2016, immediately after the 58th American Society of Hematology Annual Meeting. Rather than present a resume of the proceedings, we have elected to address some of the topical translational research and clinically relevant topics in greater detail. We address recent updates in the genetics and epigenetics of MPN, the mechanisms of transformation by mutant calreticulin, advances in the biology and therapy of systemic mastocytosis, clinical updates on JAK2 inhibitors and other therapeutic approaches for patients with MPNs, cardiovascular toxicity related to tyrosine kinase inhibitors and the concept of treatment-free remission for patients with CML.

1. Introduction

Persons with myeloproliferative neoplasms (MPNs) have been well-served by translational research over the past two decades, and in the instance of chronic myeloid leukemia (CML), for the past half-century. The recent publication of the long-term results of imatinib therapy with a median follow-up of 11 years, confirms the safety and efficacy of this first-generation tyrosine kinase-inhibitor (TKI) [1,2]. Clinical progress for other subtypes of MPNs has been more limited but there have been therapy advances for certain other MPNs [3]. For example, there are encouraging therapies for patients with systematic mastocytosis (SM)

and myeloid/lymphoid neoplasms with eosinophilia that are characterized by rearrangements of *PDGFRA*, *PDGFRB*, *FGFR1* or *PCM-JAK2* [4,5]. In contrast, in persons with *BCR/ABL1*-negative MPNs, ruxolitinib, a *JAK1* and *JAK2*-inhibitor, is the only newly-approved drug for persons with MPN-associated myelofibrosis (MF) and with polycythemia vera (PV). Most, if not all, other *JAK2*-inhibitors have been discontinued, principally due to unacceptable toxicity. Here, we address these and other important developments in CML and MPN biology and therapy discussed at the 7th John Goldman CML and MPN Colloquium which took place in Estoril, Portugal on the 15th October 2017, and the 11th Post-ASH CML and MPN workshop, which immediately

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followed the 58th American Society of Hematology meeting in San Diego, California.

2. What are the current genomic findings which inform the diagnosis, classification, prognosis and therapy-decisions in MPNs?

Genomic information has played a major role in the discovery of principal pathogenetic driver mutations in several subtypes of MPNs, beginning with the discovery of the *BCR-ABL1* fusion gene in persons with CML [2]. In contrast, the *BCR-ABL1* negative MPNs have multiple somatic mutations impacting disease biology and clinical outcome [6–12], typically in *JAK2*, *CALR* or *MPL*. These mutually exclusive alterations inform the WHO 2016 classification [13]. *JAK2*^{V617F} mutations are present in almost everyone with polycythemia vera (PV) and 50–60% of persons with essential thrombocythosis (ET) and primary myelofibrosis (PMF). Persons with PV but without *JAK2*^{V617F} often have mutations in *JAK2* exon 12 [14]. *CALR* mutations are typically absent in PV but present in 20–25% of persons with ET and PMF. Mutations in *MPL* are also rare in PV but present in about 10 percent, or less, of persons with ET or PMF. At present about 15 percent of persons with ET and PMF lack detectable mutations in *JAK2*, *MPL* and *CALR* mutations and termed ‘triple negative’. It is, however, likely that with the use of next-generation sequencing (NGS), additional mutations will be revealed [15–18]. Several other somatic mutations have been reported in persons with *JAK2*, *MPL* or *CALR* mutations including epigenetic modifiers (*ASXL1*, *TET2*, *EZH2*, *IDH1*, *IDH2*, *DNMT3A*), RNA splicing factors (*SRSF2*, *U2AF1*, *SF3B1*) and transcriptional regulators (*TP53*, *IKZF1*, *NF-E2*, *CUX1*) [19]. These mutations do not cause myelo-proliferation but influence the phenotype and prognosis. For example, mutations in *IDH1*, *IDH2*, *EZH2*, *ASXL1*, and *SRSF2* have all been shown to predict risk for leukemic transformation of patients with myelofibrosis. Moreover, mutations and deletions in *TP53* have been shown to predict leukemic transformation of all classic *BCR-ABL1* negative MPNs and promote transformation to acute myeloid leukemia in an experimental model with *JAK2*^{V617F} expression. Interestingly, mutations in *ASXL1*, *EZH2*, and *SRSF2* all appear to impinge on methylation of histone H3 lysine 27 methylation [20]. Fig. 1 depicts the principal genetic abnormalities associated with evolution of the *BCR-ABL1* negative MPNs.

Unique genetic hallmarks of other specific sub-types of MPNs are also described. For example, activating mutations in *CSF3R* occur in almost everyone with chronic neutrophilic leukemia (CNL) [21]. Mutations in *PDGFRA*, *PDGFRB* and *FGFR1*, and fusions, such as *PCM1-JAK2*, are associated in several myeloid and lymphoid neoplasms with eosinophilia [5,22,23]. In addition, the sequences of mutation acquisition may influence biology and clinical features of MPNs [24]. For example, if *JAK2*^{V617F} occurs before mutations in *TET2* or *DNMT3A* or acquired uniparental disomy (aUPD) of chromosome 14q, a person is more likely to develop PV than ET compared with the contrary [25,26]. Furthermore, an antecedent *TET2* mutation seems to prevent *JAK2*^{V617F} from up-regulating proliferation associated genes and may be associated with an increased risk of transformation to AML. Another alteration which can alter MPN phenotype are mutations in *EZH2* [16]. In contrast to *JAK2*^{V617F} mutations which may occur early or late in the development of MPNs, *CALR* mutations are thought to always be the initiating mutation in MPNs. There may also be differences in pathological signaling in the JAK-STAT pathway caused by different driver mutations. However, the precise contributions of driver and associated mutations linked to the phenotype of MPNs remains an enigma. Parenthetically, *JAK2* or *CALR* mutations and *BCR-ABL1* are reported in very rare cases of atypical myeloid neoplasms with a hybrid CML/MPN phenotype [27]. Familial MPNs are also rare and can be associated with a single-nucleotide germline variant in *TERT* added to germline mutations in the *JAK2* JH1 and JH2 domains and to the *MPL* transmembrane domain [16,28,29].

3. How do CALR mutant proteins promote transformation?

Following the seminal description of *CALR* mutations in MPNs in 2013, there has been a concerted effort to understand the molecular mechanisms by which mutant *CALR* proteins promote MPN development [11,12,30]. The fact that *JAK2*^{V617F}, *MPL*, and *CALR* mutations are mutually exclusive in MPNs suggested that transformation occurred through the same pathway. The JAK-STAT pathway had already been shown to mediate the transforming potential of *JAK2*^{V617F} mutations, where full cellular transformation required the presence of either *MPL*, *EPO*, or *CSF3* (G-CSF) receptors [31]. Notably, work carried out by Constantinescu and colleagues revealed that the presence of the thrombopoietin receptor encoded by *MPL* was essential for *CALR*-mediated transformation [30]. This has now been confirmed by several investigators [32–34]. Transformation of megakaryocytic cells by *CALR* mutants required the presence of *JAK2* protein as well as the interaction of *CALR* with *MPL*, occurring between the glycan binding site of *MPL* and the novel C-terminus of *CALR*, a domain generated by the MPN-associated mutations in *CALR* [30]. Further research showed that the charge and sequence of the novel C-terminal portion of *CALR* were essential for interaction with *MPL* and for transformation, as was the mutant *CALR*-mediated autoinhibition of the *CALR* P domain, which physiologically prevents binding of the N-terminal domain and the *MPL* receptor [32,33]. *CALR* mutant proteins by themselves induce low-level expression of megakaryocytic genes, potentially sensitizing 32D cells to *CALR*-mediated transformation, a finding also confirmed in Ba/F3 cells and human CD34⁺ cells [34–36]. Moreover, with the KDEL ER retention signal missing in *CALR* mutant proteins, *CALR* mutants showed abnormal subcellular localization [11,30,34]. Interestingly, mutations in *CALR* result in reductions in intracellular levels of *CALR* mutant protein and increased secretion of *CALR* [34,35]. While paracrine effects of secreted *CALR* mutant proteins were not uniformly observed, Dao and colleagues observed that secreted *CALR* mutant protein may indeed exert activity, enhancing inflammatory cytokine production in normal [37,38]. Further work is needed to elucidate the full spectrum of aberrant characteristics of *CALR* mutant proteins as well as the question of how these specific characteristics may be exploited therapeutically (Fig. 2).

4. Does midostaurin treatment accord molecular and clinical responses in patients with systemic mastocytosis?

Systemic mastocytosis (SM) is due to the pathologic accumulation of neoplastic mast cells in one or more extracutaneous organ(s) such as the bone marrow, spleen, liver, lymph nodes, or gastrointestinal tract [13,39]. The canonical *KIT*^{D816V} mutation is found in ~90% of SM patients and is the primary pathogenetic driver of the disease [40]. Advanced forms of SM include aggressive systemic mastocytosis (ASM), SM with an associated hematologic neoplasm (SM-AHN), and mast cell leukemia (MCL). ASM, SM-AHN, and MCL are associated with poor-prognosis [median overall survival (OS) of 3.5 years, 2 years, and ≤ 6 months, respectively], wherein morbidity and mortality often relate to progressive organ damage (so-called “C-findings”) by neoplastic mast cells [41,42]. In contrast to indolent SM, where treatment is geared to control of mast cell activation symptoms, such as flushing and diarrhea, cyto-reduction of mast cells is required to reverse organ damage in advanced disease [43]. Off-label therapies including pegylated interferon- α and cladribine, but not imatinib, have demonstrated reversion of organopathy in approximately 30–50% of patients, but data for their efficacy are primarily derived from small case series with limited follow-up and a heterogeneous mixture of indolent and advanced patients [44,45]. Midostaurin (N-benzoylstauroporine; PKC412) is a multikinase inhibitor that demonstrates activity against both wild type and D816V-mutated *KIT*, as well as *FLT3*, *PDGFR- α / β* , *PKC*, and *VEGFR2* [46,47]. In a small study of 26 patients with advanced SM, > 50% improvement or normalization of SM-related organ damage

Fig. 1. Genetic alterations and routes to leukemic transformation in *BCR-ABL1* negative classic myeloproliferative neoplasms (MPNs). Mutations in *IDH1*, *IDH2*, *EZH2*, *ASXL1*, and *SRSF2* have all been shown to predict risk for leukemic transformation of patients with myelofibrosis. Moreover, mutations and deletions in *TP53* have been shown to predict leukemic transformation of all classic *BCR-ABL1* negative MPNs and promote transformation to acute myeloid leukemia in an experimental model with *JAK2*^{V617F} expression. Interestingly, mutations in *ASXL1*, *EZH2*, and *SRSF2* all appear to impinge on methylation of histone H3 lysine 27 methylation. **(B)** The genetic alterations shown in **(A)** may occur in a cell with a pre-existing *JAK2*/*CALR*/*MPL* mutation to result in AML or could occur in a cell prior to acquisition of the *JAK2*/*CALR*/*MPL* mutation (in which case the patient may have both *CALR*/*JAK2*/*MPL* mutant MPN and *CALR*/*JAK2*/*MPL* wildtype AML). **(C)** As a final possibility, occasionally AML may be generated from a hematopoietic stem cell (HSC) unrelated to the HSC which gave rise to MPN.

was observed in 69% patients treated with midostaurin [48]. These results provided the impetus for a global, open-label, single-arm study of midostaurin in patients with ASM, MCL, and SM-AHN [49]. Among 89 evaluable patients, the overall response rate was 60%, of which 75%

were major responses (MRs; defined by complete reversion of organ damage) [40]. Responses were observed regardless of *KITD816V* status, prior therapy, or the presence of an AHN. After a median follow-up of 26 months, the median duration of response and median OS were 24.1

Fig. 2. Mechanisms of transformation by mutant calreticulin (CALR). **(A)** Binding of wildtype (wt) CALR protein to the thrombopoietin receptor (MPL) is inhibited by the P-domain of CALR (⊗). The novel C-terminus of mutant CALR protein antagonizes this inhibition (⊕), and mutant CALR protein can now bind to MPL via its N-terminal domain (⊕). [This figure was modified from Araki et al. Blood [33].] **(B)** Mutant CALR protein binds almost exclusively to MPL, thus inducing a megakaryocytic MPN phenotype such as essential thrombocythemia (ET) or primary myelofibrosis (PMF). Conversely, mutant *JAK2*^{V617F} protein may bind to either MPL or the erythropoietin receptor (EPOR) or the granulocyte-colony stimulating factor receptor (GCSFR) and thus can mediate phenotypes ranging from ET and PMF to polycythemia vera (PV). Both CALR and *JAK2*^{V617F} mutant proteins lead to activation of the JAK-STAT pathway and ultimately stimulate the proliferation of hematopoietic stem and progenitor cells. **(C)** Intracellularly, CALR mutants are subject to enhanced protein degradation. MPL binding occurs as early in the Golgi apparatus and endoplasmic reticulum (ER), and this stabilizes mutant CALR protein and leads to JAK-STAT activation. In addition, mutant CALR protein is secreted into the extracellular space where it may activate monocytes and induce inflammation.

and 28.7 months, respectively. Median OS in responders was 44.4 months vs. 15.4 months in non-responders (hazard ratio for death, 0.42; $p = 0.005$). Of the 16 patients with MCL, 8 responded, including 7 molecular remissions. The median OS was 9.4 months among all patients with MCL, and was not reached among responding MCL patients. Symptoms and quality of life were also significantly improved with midostaurin. The drug's ability to improve organ damage and symptoms may relate to its blockade of both mast cell proliferation and IgE-dependent mediator release [50]. Midostaurin was generally well tolerated with a manageable toxicity profile consisting mostly of gastrointestinal side effects and myelosuppression. Patients who experienced a response on midostaurin treatment in accordance with the modified Valent criteria or a $> 50\%$ reduction in bone marrow mast cell burden, and/or a reduction in the *KITD816V* allele burden, had an improved survival [50]. A recent study has also defined molecular correlates of response and progression on midostaurin. Patients with SM-AHN, who harbored *SRSF2*, *ASXL1*, or *RUNX1* mutations, known to adversely impact survival, had inferior response rates, median duration of midostaurin treatment, and OS, compared with those who tested negative [51]. Midostaurin was licensed in April 2017, by the FDA, for patients with ASM, SM-AHN, and MCL [4].

5. How long to continue treatment in patients with CML who have had molecularly undetectable disease for at least 2 years?

A principal goal of CML therapy with TKIs is the achievement of a stable molecular remission which accords successful discontinuation of therapy, a concept now known as 'treatment-free remission (TFR)'. Since second- and third-generation TKIs exhibit greater potency over imatinib, resulting in deeper molecular responses, they are associated with a greater proportion of patients achieving TFR, even though there is no OS superiority [52,53]. An unresolved challenge has been to determine how long to continue treatment in patients who achieve molecularly undetectable disease for at least 2 years [54]. Clearly this is a prudent issue with significant impact not only to the quality-of-life and side effects experienced by patients, but also the financial burdens that affect both patients and society. The concept of TFR has now been addressed in several studies [55–59]. The STIM1 study showed that, following a median follow-up of > 6 years, 38% of the patients who discontinued therapy have not had a molecular recurrence to date, with the rest having a relapse [55]. The Euro-SKI study enrolled 772 persons who were considered eligible to stop imatinib, dasatinib or nilotinib after a median of 91 months and a sustained MR4 or greater for at least 1 year. The molecular relapse-free survival at 12 months was 56%, with patients who had a MR4 of > 3.1 years duration, and no significant safety concerns were noted [Fig. 3(a)(b)] [60]. In patients who relapse, most, if not all, achieve molecular remissions on the re-institution of the previously discontinued TKI. Similar findings have been observed in other dasatinib- and nilotinib-treated persons in clinical trials. As illustration, the ENESTfreedom was a single arm phase 2 study which assessed the potential for TFR following first-line treatment with nilotinib, administered for 2 years followed by a 1-year consolidation [61]. The study cohort comprised of 215 patients, of which 190 could demonstrate sustained deep molecular response and enter the TFR phase of the study. 51.6% of these persons remained in molecular remission, and the median duration of nilotinib therapy in this study was relatively short at 3.6 years. At present, although nilotinib was granted a European regulatory approval in June 2017, for the inclusion of TFR, many specialists believe that, at least for the moment, it is probably best to discontinue the TKI therapy only within the framework of a clinical trial [62]. Most agree that discontinuation is feasible if close clinical and molecular monitoring is readily available [63,64]. Other factors which can be useful as positive predictive factors for TFR include a higher NK-cell count at the time of TKI discontinuation and a lower expression of the T-cell inhibitory receptor-ligand CD86 on plasmacytoid dendritic cells, deep molecular response and duration of TKI treatment [65,66].

Recent findings suggest that the age and gender of the person, as well as intrinsic psychological and cognitive variables, might also be important [67,68]. An alternative strategy for TFR involves de-escalation of the TKI doses for patients who have durable major molecular response or a deep molecular response, such as that demonstrated in the DESTINY study [69]. The observation that lower TKI doses could maintain responses in a significant proportion of the study cohort is attractive and merits further investigation. So far, however, no reliable biomarkers are available to identify patients that will not lose MMR following TKI discontinuation. The notion of achieving TFR successfully is clearly a significant milestone for the optimal management of persons with CML, most of whom have a life expectancy equivalent to that of the general population, following appropriate treatment with TKIs [70].

6. What can patients with MPNs expect from JAK2 inhibitors, and what comes next?

In contrast to most myeloid malignancies, MPNs appear to have a remarkably long natural history with many patients with ET and PV having a survival which is like that of the general population [21]. It is therefore important to avoid unnecessary treatment and treatment-related side effects, and all MPNs patients need to be assessed carefully. Ruxolitinib, a type 1 inhibitor of JAK1 and JAK2, was approved, in 2011, for the treatment of patients with intermediate- and high-risk MF, and, in 2016, for persons with PV refractory or intolerant to hydroxyurea [71–74]. These indications followed the results from randomized studies, which demonstrated the drug's ability to substantially alleviate symptoms, reduce splenomegaly, and improve blood counts in some patients, but without altering molecular response or reducing risk of transformation to AML. Longer-term data confirms a survival benefit from ruxolitinib treatment in MF, particularly for those patients in which there is histologic improvement in the bone marrow, though neither of the two randomized trials (COMFORT-1 and -2) were powered to measure differences in survival [3,75,76]. Also, ruxolitinib appears to exert no appreciable effect on the malignant clone and the discontinuation rate of the drug appears high. The drug's major effect appears to be suppression of inflammatory cytokine production, a major feature mediator of MPN-related symptoms, lending some support to the speculation that some of the survival benefit might stem from the anti-inflammatory properties which result in improving symptoms and/or splenomegaly [77]. It is of interest that the survival benefit was observed in both studies despite complete crossover and raises the possibility that the drug might indeed modify the natural history in some cases.

However, clearly not all persons with MF respond to ruxolitinib, responses are not enjoyed indefinitely, and challenges with cytopenias remain [3]. Consequently, there has been considerable interest in investigating the role of selective, potentially non-myelosuppressive type 1 JAK2 inhibitors, such as pacritinib and momelotinib [78,79]. Several other JAK2 inhibitors, such as fedratinib, have also been tested in a randomized trial in persons with MF (JAKARTA), and discontinued due to significant toxicity, including possible Wernicke's encephalopathy [80]. Pacritinib, a JAK2 and FLT3 inhibitor, has also been studied in randomized phase 3 studies (PERSIST-1 and -2), but accrual of PERSIST-2 was impeded by a FDA suspension because of concerns of safety with hemorrhage in both studies, which has now been lifted [78,81]. PERSIST-2 study includes persons with MF with a platelet count of $< 100,000/\text{dl}$, a cohort with an unmet clinical need. The study evaluated two different doses of pacritinib compared with best alternative and identified that pacritinib 200 mg twice a day was superior to either pacritinib 400 mg a day or best alternative therapy, including ruxolitinib, for endpoints of improving symptom burden and spleen response. Additionally, pacritinib resulted in improvements in red cell transfusions and tolerability, even in patients with significant thrombocytopenia. Further studies with pacritinib are planned to adequately identify the minimally effective dose of this agent for patients with MF.

Fig. 3. Results from the Euro-SKI study (ASH 2016) (a) Molecular recurrence-free survival (n = 755 of 772 persons enrolled); Euro-SKI study (ASH 2016) (b) Prognostic modeling (n = 405, imatinib);Euro-SKI study (ASH 2016).

Long term data from momelotinib, a JAK1 and JAK2 inhibitor, in MF suggest durable responses for cytopenias, as well as splenomegaly and symptoms, but neuropathy was not infrequently observed [79]. Momelotinib is thought to potentially improve anemia through its inhibition of activin A receptor type I and reduction of hepcidin production. The drug has now been tested in two phase 3 studies (SIMPLIFY-1 and -2), and though both studies confirmed the drug's credentials with respect to improving anemia, and SIMPLIFY-2 symptomatic improvement, the studies were negative and further development of the drug for MF discontinued [82]. Other JAK inhibitors in development for MF include the JAK2 inhibitor NS018 (NS Pharma, Japan), which has demonstrated modest activity and safety in phase 2 setting [83]. Type 2 JAK2 inhibitors, such as CHZ868, have also been tested in murine models, but currently clinical studies are not planned

[84]. A JAK1-selective inhibitor, INCB039110, a potent inhibitor of several keynote inflammatory cytokines, has also been studied in persons with MF and demonstrates modest activity with minimal myelo-suppression [85].

Rational ruxolitinib-based combinations, such as with pomalidomide, azacytidine, and decitabine, have now been reported [86–88]. In general, though well tolerated, they appear to add little to ruxolitinib's activity as monotherapy. Preliminary data from a dose seeking study of the PI3K δ Inhibitor TGR-1202 in combination with ruxolitinib was favorable [89]. Finally, new agents were reported in development for myelofibrosis, including PRM-151, a recombinant human pentraxin-2 that inhibits fibrocyte differentiation, and sotatercept, a novel soluble receptor fusion protein that relieves blockade of terminal erythropoiesis caused by TGF beta and has efficacy in MF-anemia [90,91]. Results

from randomized studies of long acting pegylated alpha interferons (2a and 2b) in high-risk PV and ET have confirmed their efficacy and safety, and non-inferior (but not superior) to hydroxyurea regarding the study endpoints of blood count control, prevention of thrombotic complications, and bone marrow response criteria; some ropeginterferon (2b)-treated patients also demonstrated molecular responses following one year of treatment [92–94].

7. Vascular and cardiac toxicity of ABL1-tyrosine kinase inhibitors

Following longer term follow-up, the clinical risk of ABL1-TKI related vascular and cardiac safety is increasingly recognized and has raised concerns, when TFR might not be feasible and long-term therapy is indicated [95]. The risks are greater with the second- and third-generation TKIs, and in the presence of recognized comorbidities. In general, the risks are low, but it is likely that these risks have been underestimated [96]. Several systemic reviews and meta-analysis reveal dasatinib, nilotinib, and ponatinib, but not bosutinib, to increase vascular occlusive events, compared with imatinib [97–99]. In view of these considerations, expert panels, such as the European LeukemiaNet (ELN), have convened to offer recommendations for the management and avoidance of vascular and cardiac adverse events [100]. Since the precise underlying mechanism for these effects remains an enigma, probably related to inhibition of ligand-related signaling pathways and direct effects on vascular and perivascular cells, rationale approaches to minimize the long-term sequelae from such critical adverse events have been proposed. The notion of considering the effect of aspirin and anticoagulation on TKIs induced thrombotic events in CML has been tested and found to decrease the thrombotic events excluding myocardial infarction [101]. Clearly such approaches need further study and validation, as they are crucial for the optimal management of patients with CML.

8. Conclusions

Ongoing studies in MPNs have underlined the remarkable long-term success of TKIs in the therapy of CML. This contrasts with other subtypes of MPNs where, so far, only a qualified success has been noted. With this success of TKIs in CML, the importance of TFR has been recognized as an important future milestone. This has now been achieved with the recent regulatory inclusion of nilotinib for TFR and marks the CML community's efforts to discontinue TKIs effectively, to reduce the significant therapy-related impact on the quality-of-life, other side-effects, and financial burdens that affect both patients and society, in a responsible manner. And we need rational approaches to reduce the risk of vascular and cardiac toxicities. We have also learned much from the longer-term studies of ruxolitinib's efficacy and safety and that of emerging JAK2 and other therapies for MPNs. Ruxolitinib affords substantial clinical benefits, such as symptomatic relief, reduction of splenomegaly, and a modest survival benefit. Importantly there is some uncertainty as to precise disease-initiating events in MF and efforts based on targeting multiple molecular pathways, in addition to JAK-STAT, are now assessing the role of combining ruxolitinib with other drugs, including IFN- α . Furthermore, clinical trials assessing the efficacy of the type II JAK inhibitors are now in progress. A principal challenge is to establish the prognostic and predictive impact of the numerous mutations found in *BCR-ABL1* negative classic MPN in efforts to better identify drug targets and understand the impact of the microenvironment and inflammation in the pathogenesis of these diseases.

Authorship contributions

TIM, JG, RPG, RM, SK, HJK, and OAW designed the outline strategy of the manuscript, analyzed and interpreted the data and wrote the first draft of the manuscript. All authors participated in writing significant sections of the paper and all approve the final version of the

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