

UC Irvine

UC Irvine Previously Published Works

Title

Children and Adolescents with Chronic Myeloproliferative Neoplasms: Still an Unmet Biological and Clinical Need?

Permalink

<https://escholarship.org/uc/item/4642p3w8>

Journal

HemaSphere, 3(5)

ISSN

2572-9241

Authors

Mughal, Tariq I
Deininger, Michael W
Kucine, Nicole
et al.

Publication Date

2019-10-01

DOI

10.1097/hs9.0000000000000283

Copyright Information

This work is made available under the terms of a Creative Commons Attribution License, available at <https://creativecommons.org/licenses/by/4.0/>

Peer reviewed

Children and Adolescents with Chronic Myeloproliferative Neoplasms: Still an Unmet Biological and Clinical Need?

Tariq I. Mughal¹, Michael W. Deininger², Nicole Kucine³, Giuseppe Saglio⁴, Richard A. Van Etten⁵

Correspondence: Tariq I. Mughal (e-mail: tmughal911@hotmail.com).

The remarkable story of the classic chronic myeloproliferative neoplasms (MPN), a group of clonal hematological malignancies characterized by excessive accumulation of one or more myeloid cell lineages and an inherent ability to transform to acute leukemia, and views on its molecular genetics, biology and treatment continue to evolve.^{1,2} These neoplasms comprise of chronic myeloid leukemia (CML), defined by the presence of *BCR-ABL1*, and the *BCR-ABL1*-negative disorders, polycythemia vera (PV), essential thrombocythemia (ET) and primary myelofibrosis (PMF). The term myelofibrosis (MF) includes primary myelofibrosis (PMF) and post-PV and post-ET MF. Though the diagnostic criteria and classification in accordance to the revised 2016 WHO MPN criteria are often utilized in adults, adolescents and children, there are emerging data pointing to genetic differences which impact phenotypic characteristics, treatment and prognosis.^{3,4} In addition, most of the current risk scores used for adults have not been validated in children and adolescents, with the sole exception of the EUTOS long term survival score (ELTS) that may be used to predict progression-free survival, but not overall survival (OS).^{5,6} At the thirteenth post-American Society of Hematology (ASH) Workshop on CML and MPNs, which took place on the 4th to 5th December 2018 in San Diego, some of the unique biology, emerging age-related genomic features and treatment challenges

pertaining to childhood and adolescents MPN were reviewed, with updated discussions since. Here, we briefly summarize these findings and indicate where these may have important clinical management implications; broader aspects of the workshop are published elsewhere.⁷

Epidemiology and genetics

With the sole exception of juvenile myelomonocytic leukemia (JMML) and possibly cutaneous mastocytosis, the incidence of all MPNs increases with age.⁸ This observation lends support to the potential link between aging and inflammation ('inflammaging') as a driver of clonal evolution and selection in MPNs.⁹ MPNs are extremely rare in children and adolescents, and even rarer in infants. Also, unlike in adult MPNs, clonal hematopoiesis of indeterminate potential (CHIP) is unlikely in pediatric populations, and hence unlikely to play a role in the pathogenesis of pediatric MPNs.^{10,11} That said, there is some evidence suggestive of the notion of colonization of hematopoietic tissues by intestinal microbiota being associated with the development of MPN in *TET2* deficient mice, a process mediated by IL-6 and linking progression from CHIP to MPNs to extrinsic 'macro-environmental' factors.¹²⁻¹⁴ Such observations lend some support to the hypothesis supporting the risk of a history of infection or autoimmune disease and myeloid malignancies.¹⁵

Children and adolescents with CML do not appear to have cytogenetic abnormalities in addition to the Philadelphia chromosome, but have been observed to harbor different *BCR-ABL1* breakpoint distribution and additional somatic mutations, often resembling those seen in *BCR-ABL1*-positive acute lymphoblastic leukemia (ALL).^{16,17} The leukemogenesis of pediatric *BCR-ABL1*-negative MPNs is associated with genetic drivers that are similar to those in adults, but with important differences. For example, a lower frequency of mutations in all 3 founding driver mutations, *JAK2*, *MPL* and *CALR*, and an absence of mutations in any of the about 100 known MPN-associated genes have been reported in a significant proportion of children and adolescents diagnosed with MPNs.¹⁸⁻²⁰ It is of interest that many children without these mutations have a diagnosis of hereditary thrombocytosis or ET. Some of these children were reported to harbor *MPL*^{S505A}, which is actually *MPL*^{S505N}, and in keeping with the original report in 2007.^{21,22} The precise significance and role of co-existing germline, driver and secondary somatic mutations in relation to age at the onset of the MPN clone are also poorly understood.²³ Younger patients appear to develop PV

Conflicts of interest Disclosures

Deininger MW: No relevant disclosures.

Kucine N: No relevant disclosures.

Mughal TI: Consultancy: Roche/FMI, Stemline, Sangamo; Speakers' Bureau: Sanofi.

Saglio G: Consultancy: Bristol-Myers Squibb.

Research funding: Novartis.

Van Etten RA: Scientific Advisory Boards: Bristol Myers-Squibb, Pfizer, TEVA, Sunesis.

¹Tufts University Cancer Center, Boston, MA, USA

²University of Utah, Salt Lake City, UT, USA

³Weill Cornell Medicine, New York, NY

⁴Orbassano University Hospital, Turin, Italy

⁵University of California Irvine, Irvine, CA, USA.

Copyright © 2019 the Author(s). Published by Wolters Kluwer Health, Inc. on behalf of the European Hematology Association. This is an open access article distributed under the Creative Commons Attribution License 4.0 (CCBY), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

HemaSphere (2019) 3:5(e283)

Received: 21 June 2019 / Accepted: 26 June 2019

Citation: Mughal TI, Deininger MW, Kucine N, Saglio G, Van Etten RA. Children and Adolescents with Chronic Myeloproliferative Neoplasms: Still an Unmet Biological and Clinical Need?. *HemaSphere*, 2019;3:5. <http://dx.doi.org/10.1097/HS9.0000000000000283>.

following the acquisition of a single somatic *JAK2*^{V617F} mutation, in contrast to older patients in whom this mutation is often accompanied by secondary mutations.²⁴ Interestingly, a diagnosis of MPN has been made in several children who were previously observed to have *JAK2*^{V617F} mutations detected in the newborn screening cards. Indeed, data from the Swedish-familial cancer registry suggest a genetic predisposition to acquiring MPN driver mutations at younger ages in patients with a high-risk family history. It is likely that inherited and environmental factors are shared in this genetic predisposition.²⁵ Indeed, anecdotal reports of a *TERT* single-nucleotide variant has also been reported, especially in females, and associated with exposure to benzene and toluene.²⁶

The International Pediatric Registry data suggest the incidence of CML in children and adolescents to be considerably lower than that in adults, with an annual incidence of about 0.7 per 1,000,000 and 1.2 per 1,000,000, respectively.²⁷ The incidence of PV and ET has been estimated to be about 1 to 2 cases per 10,000,000 children and adolescents annually; the incidence of PMF and MF is unknown and considered extremely low.²⁸ There are also some reports suggesting that with increased pediatric-MPN awareness, incidence rates could rise.

Clinical and treatment issues in pediatric MPNs

Clinically, children and adolescents with CML in chronic phase often present with proliferative features, such as splenomegaly and higher leucocyte counts, compared with adults. Additionally, a greater proportion of them have advanced disease at diagnosis.²⁹ Pediatric *BCR-ABL1*-negative MPN patients can present with thrombotic events, in particular when associated with *JAK2* mutations.^{30–32} The risk of all thrombosis is quite low overall, estimated at about 9% for PV and 4% for ET. In contrast, the incidence of venous thrombotic events appears to be quite high – estimated at about 85% in the London series, whilst hemorrhagic complications are rare (<5%).³² Children with PV and ET also tend to exhibit higher white blood counts and often present with hepato-splenomegaly.³²

For adult patients with CML, the licensed *ABL1*-tyrosine kinase inhibitors (TKI), accord an OS not dissimilar from that of the general population, and about one-half of patients achieving sustained deep molecular remission (DMR) are able to discontinue therapy successfully and durably.¹ In contrast, though 3 TKIs are now approved for the treatment of CML in children and adolescent, the data with regards to efficacy, safety and treatment-discontinuation is considerably less robust, and uncertainty with regards to the optimal treatment remains.⁵ At present imatinib, nilotinib, and dasatinib are approved for first-line therapy. Imatinib has clearly improved clinical outcomes, and both nilotinib and dasatinib appear to achieve faster and deeper molecular responses, similar to those seen in adults. It is, however, unclear as to which TKI to offer initially, and no randomized studies comparing imatinib to nilotinib or dasatinib have been conducted so far. Neither are there any specific ELN or NCCN guidelines for treatment or monitoring of children and adolescents with CML; rather the adult guidelines are often extrapolated.²⁷ Compliance and adherence might be a bigger issue in adolescents and requires careful monitoring.

There is also considerable concern with regards to the long-term side effects of TKIs being administered during extended periods during growth and development, implying that treatment free remission (TFR) is an even more important goal than in adults.

Further, the notion of TFR remains largely untested in pediatric populations, creating challenges in developing optimal treatment algorithms for children and adults with CML in chronic phase. A recent international study, STOP IMAPED, assessed imatinib discontinuation in 14 children with CML in chronic phase who had achieved and maintained DMR for >2 years.³³ The study observed that the majority of study cohort experienced a molecular relapse following imatinib discontinuation with an overall probability of maintaining DMR at 6 months being 28.6%. In contrast to the recent EURO-SKI adult study, STOP IMAPED failed to reveal any predictive parameters, such as age, sex, time from diagnosis to the onset of therapy, treatment duration or duration of DMR until discontinuation, to be associated with molecular relapse.³⁴ The precise reasons for these discordant results is unclear. It can be hypothesized that the presence of more proliferative features at diagnosis, different *BCR-ABL1* breakpoint distribution, and possible differences in the immune response between children and adults have contributed. Indeed, at present we know very little about the immunological mechanisms which may be relevant in maintaining DMR following TKI discontinuation. Regardless, further prospective and larger studies assessing discontinuing TKIs successfully are urgently required in children and adolescents with CML prior to the introduction of TFR into the pediatric clinics outside of clinical trials.

The recent approval of nilotinib and dasatinib for pediatric patients has also complicated the decision-making process regarding allogeneic stem transplantation (allo-SCT). Clearly, patients who have a suboptimal response or failure to 2 lines of TKIs should be considered for an allo-SCT. Allo-SCT should also be offered to CML patients who acquire the T315I mutation in *BCR-ABL1*, since the third generation TKI, ponatinib, is not currently licensed for children and adolescents. A recent comprehensive survey of 247 US-based pediatric hematologists' attitude towards an optimal treatment algorithm suggest that most do not support the use of allo-SCT for first-line therapy, and about 40% support it if suitable donor was available.³⁵ However, the vast majority support the use of allo-SCT following 2 lines of TKI therapy.

For adult patients with MF a qualified therapeutic success with a substantial relief of disease-related symptoms, splenomegaly and survival benefit has been observed with the only licensed *JAK1/2* inhibitor, ruxolitinib,^{36,37} but therapy-related anemia is often an anticipated downside.³⁸ Allo-SCT, however, remains the only treatment which can achieve long term remission and potential cure, but is associated with considerable morbidity and mortality.³⁹ There are, however, scanty data with regards to the use of *JAK* inhibitors and allo-SCT in children and adolescents. Indeed, at present, there is no firm consensus on the optimal clinical management of children with PV, ET, or MF. Hydroxyurea is used frequently, but long-term safety remains unclear, and there is limited data available on use of interferon alpha in this patient population.^{30,32}

Future prospects

Arguably, despite much progress in the field of MPNs, and in particular CML, many important issues pertaining to the diagnosis, classification, risk stratification and treatment of children and adolescents remain. Mutational profiling has revealed important differences between adult and pediatric MPNs which influence the biology and prognosis, but larger contemporary cohorts with long-term follow-up are needed to assess this and the impact on treatment appropriately. In this

regard, it is of note that during this year's International Childhood Cancer Awareness Day, both the Society for Pediatric Oncology Europe (SIOPE) and Childhood Cancer International (CCI) on Pediatric Hematology and Oncology launched a manifesto calling for the European Parliament to maintain and reinforce support. The rarity of pediatric MPNs and the increasing recognition of long-term side-effects of successful treatments, including cardiovascular, endocrine, growth and psychological issues underscore the urgent need for agreeable funding and collaborations worldwide.

Acknowledgments

The authors thank the participants and faculty of the thirteenth Post-ASH CML-MPN workshop, Dr Alpa Parmar for her organizational skills, and Incyte Corporation, Novartis Oncology, Pharma Essentia and Alpine Oncology Foundation for unrestricted grant support.

References

- Mughal TI, Radich JP, Deininger MW, et al. Chronic Myeloid Leukemia: Reminiscences and dreams. *Haematologica*. 2016;101:541–558.
- Deininger MW, Tyner JW, Solary E. Turning the tide in myelodysplastic/myeloproliferative neoplasms. *Nat Rev Cancer*. 2017;17:425–440.
- Arber DA, Orazi A, Hasserjian R, et al. The 2016 revision to the World Health Organization classification of myeloid neoplasms and acute leukemia. *Blood*. 2016;127:2391–2405.
- Wong WJ, Hasserjian R, Pinkus GS, et al. JAK2, CALR, MPL and ASXL1 mutational status correlates with distinct histological features in Philadelphia chromosome-negative myeloproliferative neoplasms. *Haematologica*. 2018;103:e63–e68.
- Hijiya N, Suttrop M. *How I treat chronic myeloid leukemia in children and adolescents blood*. 2019;pre-published, doi: 10.1182/blood.2018882233.
- Gurrea-Salas D, Glauche I, Tauer JT, et al. Can prognostic scoring systems for chronic myeloid leukemia as established in adults be applied to pediatric patients? *Ann Hematol*. 2015;94:1363–1371.
- Mughal TI, Pemmaraju N, Radich JP, et al. Emerging translational science discoveries, clonal approaches and treatment trends in chronic myeloproliferative neoplasms. *Hematol Oncol*. 2019;1–13. doi: 10.1002/hon.2622.
- Srouf SA, Devesa SS, Morton LM, et al. Incidence and patient survival of myeloproliferative neoplasms and myelodysplastic/myeloproliferative neoplasms in the United States, 2001–12. *Br J Haematol*. 2016;174:382–396.
- Fulop T, Witkowski JM, Olivier F, et al. The integration of inflammatory in age-related diseases. *Semin Immunol*. 2018;40:17–35.
- Elias HK, Bryder D, Park CY. Molecular mechanisms underlying lineage bias in aging hematopoiesis. *Semin Hematol*. 2017;54:4–11.
- Chambers SM, Shaw CA, Gatz C, et al. Aging hematopoietic stem cells decline in function and exhibit epigenetic dysregulation. *PLoS Biol*. 2007;5:e201.
- Jaiswal S, Fontanillas P, Flannick J, et al. Age-related clonal hematopoiesis associated with adverse outcomes. *N Engl J Med*. 2014;371:2488–2498.
- Fleischman AG, Aichberger KJ, Luty SB, et al. TNF alpha facilitates clonal expansion of JAK2V617F positive cells in myeloproliferative neoplasms. *Blood*. 2011;118:6392–6398.
- Meisel M, Hinterleitner R, Pacis A, et al. Microbial signals drive pre-leukaemic myeloproliferation in a Tet2-deficient host. *Nature*. 2018;557:580–584.
- Kristinsson SY, Björkholm M, Hultcrantz M, et al. Chronic immune stimulation might act as a trigger for the development of acute myeloid leukemia or myelodysplastic syndromes. *J Clin Oncol*. 2011;29:2897–2903.
- Hijiya N, Schultz KR, Metzler M, et al. Pediatric chronic myeloid leukemia is a unique disease that requires a different approach. *Blood*. 2016;127:392–399.
- Krumbholz M, Karl M, Tauer JT, et al. Genomic BCR-ABL1 breakpoints in pediatric chronic myeloid leukemia. *Genes Chromosomes Cancer*. 2012;51:1045–1053.
- Karow A, Nienhold R, Lundberg P, et al. Mutational profile of childhood myeloproliferative neoplasms. *Leukemia*. 2015;29:2407–2409.
- Ernst T, Busch M, Rinke J, et al. Frequent ASXL1 mutations in children and young adults with chronic myeloid leukemia. *Leukemia*. 2018;32:2046–2049.
- Kucine N, Viny AD, Rampal R, et al. Genetic analysis of five children with essential thrombocythemia identified mutations in cancer-associated genes with roles in transcriptional regulation. *Haematologica*. 2016;101:e237–e239.
- Defour J-P, Levy G, Leroy E, et al. The S505A thrombopoietin receptor mutation in childhood hereditary thrombocythemia and essential thrombocythemia is S505N: single letter amino acid code matters. *Leukemia*. 2019;33:563–564.
- Teofili L, Giona F, Martini M, et al. Markers of myeloproliferative diseases in childhood polycythemia vera and essential thrombocythemia. *J Clin Oncol*. 2007;25:1048–1053.
- Kelly K, McMahon C, Langabeer S, et al. Congenital JAK2V617F polycythemia vera: where does the genotype-phenotype diversity end? *Blood*. 2008;112:4356–4357.
- Fowles JS, How J, Allen MJ, et al. Young versus old age at diagnosis confers distinct genomic profiles in patients with polycythemia vera. *Leukemia*. 2019;Pre-pub April 2019; doi.org/10.1038/s41375-018-0349-9.
- Sud A, Chattopadhyay S, Thomsen H, et al. Familial risk of acute myeloid leukemia, myelodysplastic syndromes, and myeloproliferative neoplasms. *Blood*. 2018;132:973–976.
- Trifa AP, Bănescu C, Tevet M, et al. TERT rs2736100 A>C SNP and JAK2 46/1 haplotype significantly contribute to the occurrence of JAK2 V617F and CALR mutated myeloproliferative neoplasms – a multicentric study on 529 patients. *Brit J Haem*. 2016;174:218–226.
- Gunes AM, Millot F, Kalwak K, et al. Features and Outcome of Chronic Myeloid Leukemia (CML) at Very Young Age: Data from the International Pediatric CML Registry (I-CML-Ped Study). *Blood*. 2018;132:1748.
- Hofmann I. Myeloproliferative neoplasms in children. *J Hematop*. 2015;8:143–157.
- Millot F, Guilhot J, Suttrop M, et al. Advanced phases at diagnosis of childhood chronic myeloid leukemia: the experience of the international registry for chronic myeloid leukemia (CML) in children and adolescents (I-CML-Ped Study). *Blood*. 2017;130:316.
- Barbui T. How to manage children and young adults with myeloproliferative neoplasms. *Leukemia*. 2012;26:1452–1457.
- Kucine N, Al-Kawaaz M, Hajje D, et al. Difficulty distinguishing essential thrombocythemia from polycythemia vera in children with JAK2V617F-positive myeloproliferative neoplasms. *Br J Haematol*. 2019;185:136–139.
- Ianotto JC, Curto-Garcia N, Lauermannova M, et al. Characteristics and outcomes of patients with essential thrombocythemia or polycythemia vera diagnosed before 20 years of age, a systematic review. *Haematologica*. 2019;epub ahead of print, doi: 10.3324/haematol.2018.200832.
- Brujin C, Millot F, Suttrop M, et al. Discontinuation of imatinib in children with chronic myeloid leukaemia in sustained deep molecular remission: results of the STOP IMAPED study. *Br J Haem*. 2019;185:718–724.
- Sauselle S, Richter J, Guilhot J, et al. Discontinuation of tyrosine kinase inhibitor therapy in chronic myeloid leukemia (EURO-SKI): a prespecified interim analysis of a prospective, multicentre, non-randomised, trial. *Lancet Oncol*. 2018;19:747–757.
- Andolina JR, Burke MJ, Hijiya N, et al. Practice patterns of physician treatment for pediatric chronic myelogenous leukemia. *Biol Blood Marrow Transplant*. 2019;25:321–327.
- Verstovsek S, Mesa RA, Gotlib J, et al. Long-term treatment with ruxolitinib for patients with myelofibrosis: 5-year update from the randomized, double-blind, placebo-controlled, phase 3 COMFORT-I trial. *J Hematol Oncol*. 2017;10:55.
- Harrison CN, Vannucchi AM, Kiladjan JJ, et al. Long-term findings from COMFORT-II, a phase 3 study of ruxolitinib vs best available therapy for myelofibrosis. *Leukemia*. 2016;30:1701–1707.
- Naymagon L, Mascarenhas J. Myelofibrosis-related anemia: current and emerging therapeutic strategies. *HemaSphere*. 2017;1:e1.
- Passamonti F. Stem cell transplant in MF: it's time to personalize. *Blood*. 2019;133:2118–2120.