Illuminating the mutational spectrum of pediatric myeloproliferative neoplasms

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The myeloproliferative neoplasms (MPN) are clonal, hematopoietic stem cell-derived diseases characterised by bone marrow proliferation of one or more of the myeloid cell lineages with the main subtypes being polycythemia vera (PV), essential thrombocythemia (ET), and primary myelofibrosis (PMF). The diagnosis and classification of these diseases is dependent on clinical, hematological, histo-morphological and increasingly, on molecular genetic findings [1]. In adults, the most commonly acquired mutation associated with MPN is the JAK2 V617F, occurring in >95% of PV cases and in 50 - 60% of ET and PMF patients. This mutation disrupts the function of the intracellular JAK2 molecule that is required for normal hematopoietic cytokine signalling and has become a target for therapeutic intervention [2]. Alternative mutations of JAK2 within exon 12 can be detected in the majority of those remaining V617F-negative PV patients [3]. Several other genetic abnormalities have been described in MPN which occur at a much lower frequency than the JAK2 V617F. Mutations in MPL exon 10, which encodes the receptor for thrombopoietin, a major regulator of megakaryocyte and platelet development, are present in approximately 5% and 3% of PMF and ET patients respectively [4]. Recent whole exome sequencing studies of adult MPN patients have led to the discovery of insertion and/or deletion mutations in exon 9 of CALR, a gene that encodes the endoplasmic reticulum-associated, calcium binding protein, calreticulin. These disease driving mutations of CALR result in a frame shift of the coding sequence, altered amino acid composition of the translated protein, and loss of the endoplasmic reticulum retention motif. CALR mutations do not occur in PV patients but are present in up to 80% of ET and PMF patients who are JAK2 V617F-and MPL-mutation negative [5]. Approximately 10% of adult ET and PMF patients have no evidence of these three driver mutation types and are termed “triple-negative” MPN. Supplementary “passenger” mutations may be acquired that can influence the initiation and evolution of the disease, however these mutations are not specific for MPN as they are also present in other myeloid malignancies such as myelodysplastic syndromes and acute myeloid leukemia [6].

Sporadic MPN are uncommon in children with an incidence approximately one hundred fold lower than that of adults suggesting an alternative aetiology of the disease in such cases [7,8]. In childhood MPN, the prevalence of the JAK2 V617F mutation is lower when compared to that in adults [9-11] with acquired mutations of MPL exceedingly rare [12]. Analysis of the more recently described CALR mutations in pediatric MPN cohorts has also revealed significantly lower mutation rates than in adult MPN, inferring differences in the molecular pathogenesis of these diseases in childhood [13,14]. Furthermore, the relatively high proportion of pediatric cases without evidence of one the three driver mutation types associated with adult MPN suggests such cases may not be clonal malignancies: this has significant implications for therapeutic intervention with cytoreductive agents [15].

Sheding further light on this issue, several groups have recently applied targeted sequencing of large panels of either myeloid- or cancer-associated genes to pediatric MPN, especially ET patients [16-18] with each of these contemporary studies revealing insights into the mutational spectrum with the results summarized in Table 1. Karow et al. [16] analysed 43 pediatric MPN patients, most of whom had ET, by using capture-based targeted next-generation sequencing to simultaneously search for mutations in 104 hematopoietic malignancy-associated genes. The JAK2 V617F and CALR mutations were found most frequently, however the frequency of mutations in genes implicated in epigenetic regulation (ASXL1, DNMT3A, EZH2, IDH1, and TET2) was significantly lower than that of 25% observed in adult MPN. Confirming previous studies, 30% of patients had no evidence of mutations in MPN-associated genes. A second study of 25 childhood ET cases employed targeted sequencing of 55 genes associated with myeloid malignancy [17]. Again JAK2 and CALR mutations were less frequent than in adult cohorts but somewhat conversely, ASXL1 mutations were demonstrated in four cases (Table 1). Lastly, using a targeted deep sequencing assay of 585 cancer-related genes in five characterized childhood ET patients, Kucine et al. [18] identified mutations, in addition to the JAK2 V617F, in genes involved in transcriptional regulation (NUP98, MED12, AR, and CEBPA) (Table 1). Noticeably, each of these examinations has identified mutated co-operating genes but with minimal inter-study cross-over, possibly representing differences in methodological approach and the size of the gene panels utilised.

An interesting observation from these latter three studies is the high rate of co-occurrence of mutations in pediatric ET which suggests that children may have a more complex disease than adults thus reflecting an unstable genetic background: rapid acquisition of these mutations may account for the early onset of ET in children. While these approaches are highly informative and have the capability to identify targets that may be clinically actionable, these exon sequencing approaches may fail to identify abnormalities in non-coding regions. Sequencing of the 5’ untranslated region of THPO, the gene that encodes the hormone

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approach for a molecular genetic diagnosis. A JAK2, MPL, or CALR mutation as a minor criterion for diagnosis of MPN is likely to include the presence of a clonal marker other than a major role [20]. The next World Health Organization classification that the pathogenesis of MPN is not entirely dependent on genetic however these investigations will require confirmation in extended investigations. Thrombopoietin which controls platelet production, has recently been shown to be of diagnostic value [19].

| Gene | Mutation frequency |
|------|--------------------|
| JAK2 V617F | 8 (19.5%) - - - 6 (24.0%) - - - 3 (60.0%) - - |
| CALR | - - - 1 (4.0%) - - - - - |
| MPL | - - - - - - - - |
| JAK2 | 4 (9.8%) - - - - - - |
| MPL | 4 (9.8%) - - - - - - |
| NUP98 | - - - - - - - - |
| CUX1 | 2 (4.9%) - - - - - - |
| DNM3A | 2 (4.9%) - - - - - - |
| EF300 | 2 (4.9%) - - - - - - |
| EPO | 2 (4.9%) - - - - - - |
| MYB | 2 (4.9%) - - - - - - |
| NF1 | 2 (4.9%) - - - - - - |
| PTPRT | 2 (4.9%) - - - - - - |
| MED12 | - - - - - - - - |
| DAX | - - - - - - - - |
| AR2 | - - - - - - - - |
| AKT3 | 1 (2.4%) - - - - - - |
| FOXP1 | 1 (2.4%) - - - - - - |
| IDH1 | 1 (2.4%) - - - - - - |
| IDH3 | 1 (2.4%) - - - - - - |
| JAK2 | 1 (2.4%) - - - - - - |
| PIK3AP1 | 1 (2.4%) - - - - - - |
| PIK3C2A | 1 (2.4%) - - - - - - |
| RBL2 | 1 (2.4%) - - - - - - |
| SH2B3 | 1 (2.4%) - - - - - - |
| FLT3 | - - - - - - - - |
| U2AF1 | - - - - - - - - |
| NTR | - - - - - - - - |
| MLT | - - - - - - - - |
| GNAS | - - - - - - - - |
| RUNX1 | - - - - - - - - |
| WT1 | - - - - - - - - |
| ZRS2 | - - - - - - - - |
| STAF5B | - - - - - - - - |
| CEBPA | - - - - - - - - |
| TET1 | - - - - - - - - |
| TET9 | - - - - - - - - |
| CEBPA | - - - - - - - - |
| ZRS2 | - - - - - - - - |
| STAF5B | - - - - - - - - |
| CEBPA | - - - - - - - - |
| TET1 | - - - - - - - - |
| TET2 | - - - - - - - - |
| FLT3 | - - - - - - - - |
| U2AF1 | - - - - - - - - |
| NRT | - - - - - - - - |
| MIL | - - - - - - - - |
| GNAS | - - - - - - - - |
| RUNX1 | - - - - - - - - |
| WT1 | - - - - - - - - |
| ZRS2 | - - - - - - - - |
| STAF5B | - - - - - - - - |
| CEBPA | - - - - - - - - |
| TET1 | - - - - - - - - |
| TET2 | - - - - - - - - |

In conclusion, targeted sequencing of pediatric MPN patients has revealed some of the underlying complexity of co-operating mutations however these investigations will require confirmation in extended cohorts with functional validation of mutations. It must be remembered that the pathogenesis of MPN is not entirely dependent on genetic abnormalities but other causes, such as inflammation, are likely to play a major role [20]. The next World Health Organization classification of MPN is likely to include the presence of a clonal marker other than a JAK2, MPL, or CALR mutation as a minor criterion for diagnosis of ET or PMF [21], expediting the adoption of a targeted exon sequencing approach for a molecular genetic diagnosis.

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