Methylation of miR-34a, miR-34b/c, miR-124-1 and miR-203 in Ph-negative myeloproliferative neoplasms

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Abstract
Background: MicroRNA (miR) miR-34a, -34b/c, -124-1 and -203 are tumor suppressor miRs implicated in carcinogenesis.

Methods: We studied DNA methylation of these miRs in Philadelphia-negative (Ph-ve) myeloproliferative neoplasms (MPNs). Methylation-specific PCR (MSP), verified by direct sequencing of the methylated MSP products, was performed in cell lines, normal controls and diagnostic marrow samples of patients with MPNs.

Results: Methylation of these miRs was absent in the normal controls. miR-34b/c were homozygously methylated in HEL cells but heterozygously in MEG-01. In HEL cells, homozygous miR-34b/c methylation was associated with miR silencing, and 5-aza-2′-deoxycytidine treatment led to re-expression of both miR-34b and miR-34c, consistent with that both miRs are under the regulation of the same promoter CpG island. miR-34a was heterozygously methylated in MEG-01 and K-562, miR-203 was completely unmethylated in K-562 and SET-2 but no MSP amplification was found in both HEL and MEG-01, suggestive of miR deletion. In primary samples, four each had miR-34b/c and -203 methylation, in which two had concomitant methylation of miR-34b/c and -203. miR-34a was methylated in one patient and none had methylation of miR-124-1. Four patients (15.6%) had methylation of at least one of the four miRs. miR methylation did not correlate with clinical parameters, disease complications or JAK2 V617F mutation.

Conclusion: This is the first report of miR hypermethylation in MPNs. miR-203 hypermethylation is not specific to Ph+ve leukemias but also present in Ph-ve MPNs. miR-34b/c methylation was associated with reversible miR silencing. There was no correlation of miR methylation with clinical demographic data or outcome.

Keywords: microRNA, tumor suppressor, hypermethylation, Ph-negative myeloproliferative neoplasm

Background
Philadelphia-negative (Ph-ve) myeloproliferative neoplasm (MPN) is a stem cell disease with proliferation of myeloid lineage, leading to the development of distinct clinical entities including polycythemia vera (PV), essential thrombocytopenia (ET) and primary myelofibrosis (PMF) [1-3]. JAK2 V617F mutation, resulting in constitutive activation of JAK-STAT signaling, occurs in about half of the patients with ET and PMF but in more than 90% of patients with PV [1]. Gene methylation is an alternative mechanism of gene inactivation, and various tumor suppressor genes regulating the cell cycle, apoptosis and cell signaling have been shown to be hypermethylated in hematological malignancies [4].

MicroRNA (miR) is a single-stranded, non-coding RNA molecule of 22-25 nucleotides, which leads to downregulation of target protein expression [5]. miRs are involved in carcinogenesis [6]. miRs can be either oncogenic (oncomir) when tumor suppressor genes (TSG) are targeted, or tumor suppressive (tumor suppressor miRs) when oncogenes are targeted [7].

Recently, miR-34a, miR-34b/c, miR-124-1 and miR-203 hypermethylation have been implicated in carcinogenesis.
Hypermethylation of miR-34a, a transcriptional target of p53, has been demonstrated in solid and hematopoietic cancers [8,9], whereas restoration of which will inhibit CDK6 translation by complementary binding to the 3’ untranslated region (3’ UTR) of the CDK6 mRNA and induce apoptosis, thereby showing the tumor suppressor role of miR-34a [8]. Epigenetic inactivation of miR-34b, another p53 downstream target of the miR-34 family, has also been implicated in acute myeloid leukemia (AML), and the re-expression of miR-34b led to suppression of CREB expression and inhibition of cell proliferation [10]. Promoter methylation of miR-124-1, the first tumor suppressor miR found to be regulated by DNA methylation, has been shown to confer poor prognosis of acute lymphoblastic leukemia (ALL) [11]. Moreover, hypermethylation of miR-203 has been reported in chronic myeloid leukemia (CML), conferring a proliferative advantage to the tumor cells by inhibiting the oncogenic BCR-ABL fusion protein [12]. In Ph-ve MPN, little is known about the epigenetic alteration of miR methylation. In this report, we studied the methylation status of miR-34a, miR-34b/c, miR-124-1 and miR-203 in PV, ET and PMF.

Methods

Patient samples
DNA was extracted from primary marrow samples at diagnosis of 45 patients with MPN [ET, N = 34 (75.5%); PV, N = 8 (17.8%) and PMF, N = 3 (6.7%)]. There were 24 (53.3%) male and 21 (46.7%) female patients with a median age of 67.5 years (range: 28 - 89 years), a median presenting platelet count of 848 × 109/L (range: 196 - 2275 × 109/L), a median presenting hemoglobin level (Hb) 13.3 g/dL (range: 9-22 g/dL), and a median presenting leukocyte count of 14.4 × 109/L (range: 7-28 × 109/L). Apart from five (11.1%) patients in whom the presenting hematological examination at presentation were not available for review, 25 (62.5%) were asymptomatic at diagnosis, four (10%) with bleeding, four (10%) with erythromelalgia, two (5%) with minor stroke, three (7.5%) with abdominal pain, and one each (2.5%) with blurred vision and weight loss. Of 39 patients with adequate follow-up information, five (12.8%) had myeloid transformations (MDS or AML) at the time of study. Of 40 patients with data on thrombosis, nine (22.5%) had thrombotic events. Apart from 5 patients with unknown JAK2 mutation status, 26 (65%) had JAK2 V617F mutation. (Table 1) The study has been approved by Institutional Review Board of Queen Mary Hospital with written informed consent.

Cell lines and culture
MEG-01 and K-562 cells were kindly provided by Dr Mo Yang, Department of Paediatrics, Queen Mary Hospital, The University of Hong Kong, Hong Kong. HEL cells were obtained from Dr Dong-Er Zhang, Department of Pathology and Molecular Biology, Moores Cancer Center, University of California San Diego, USA. SET-2 cells were purchased from Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH (DMSZ) (Braunschweig, Germany). SET-2 was derived from ET at megakaryoblastic leukemic transformation. HEL was derived from AML M6. Both SET-2 and HEL cells carry JAK2 V617F mutation. MEG-01 and K-562 were derived from blast transformation of patients with CML. Cell cultures were maintained in RPMI media 1640 (Invitrogen, Carlsbad, CA), supplemented with 10% (20% for SET-2) fetal bovine serum (Invitrogen, Carlsbad, CA), 50 μg/ml streptomycin (Invitrogen, Carlsbad, CA), and 50 μg/ml penicillin, and in a humidified atmosphere of 5% CO2 at 37°C.

Methylation-specific polymerase chain reaction (MSP)
DNA was extracted from bone marrow samples at diagnosis and from cell lines by standard method. MSP for aberrant gene promoter methylation was performed as previously described [13,14]. Treatment of DNA with bisulfite for conversion of unmethylated cytosine to uracil (but unaffected methylated cytosine) was performed with a commercially available kit (EpiTect Bisulfite Kit, Qiagen, Germany). Primers used for the methylated MSP (M-MSP) and unmethylated MSP (U-MSP) were shown in Table 2. DNA from normal bone marrow donors was used as negative control, while enzymatically methylated control DNA (CpGenome Universal Methylated DNA, Chemicon) was used as positive control in all the experiments. MSP was performed in a thermal cycler (9700, Applied Biosystems, Foster City, CA) with the following cycling conditions: 95°C for 5 minutes, specific cycles of 95°C for 30 seconds, specific annealing temperature for 30 seconds (Table 2), 72°C for 30 seconds, and a final extension of 10 minutes at 72°C. The MSP mixture contained 50 ng of bisulfite-treated DNA, 0.2 mM dNTPs, MgCl2 (Table 2), 10 pmol of each primer, 1 × PCR buffer, and 2.5 units of AmpliTaq Gold DNA Polymerase (Applied Biosciences, Foster City, CA) in a final volume of 25 μl. Ten microliters of PCR products were loaded onto 6% non-denaturing polyacrylamide gels, electrophoresed, and visualized under ultraviolet light after staining with ethidium bromide.

5-aza-2′-deoxycytidine (5-AzadC) treatment
HEL cells were homozygously methylated for miR-34b/c. Cells were seeded in six-well plates at a density of 1 × 10^6 cells/ml, and cultured with 1.5 μM of 5-AzadC for 7 days. Cells on day 0 and day 7 of treatment were harvested.

RNA isolation and stem-loop reverse transcription-polymerase chain reaction (RT-PCR)
Total RNA was isolated using mirVana™ miRNA Isolation Kit (Ambion, Austin, TX), according to the manufacturer’s
Table 1 Patient demographic data and status of microRNA methylation

| sex | age | Diagnosis | Symptoms at diagnosis | myeloid transformation | JAK2 V617F mutation | miR-34b/c | miR-34a | miR-203 | miR-124-1 |
|-----|-----|-----------|----------------------|----------------------|---------------------|---------|--------|--------|-----------|
| F   | 78  | ET        | Epigastric pain      | No                   | yes                 | U       | U      | M      | U         |
| F   | 81  | ET        | Epigastric pain      | No                   | yes                 | U       | U      | U      | U         |
| M   | 32  | ET        | erythromelalgia      | No                   | yes                 | U       | U      | U      | U         |
| M   | 81  | ET        | N/A                  | No                   | yes                 | U       | U      | U      | U         |
| M   | 59  | ET        | N/A (No record)      | Yes                  |                     | M       | U      | U      | U         |
| M   | 76  | ET        | Nil                  | No                   | yes                 | U       | M      | U      | U         |
| M   | 53  | ET        | Nil                  | MDS                  | yes                 | U       | U      | U      | U         |
| F   | 89  | ET        | Nil                  | N/A (No record)      | yes                 | U       | U      | U      | U         |
| F   | 43  | ET        | Nil                  | N/A (No record)      | yes                 | M       | U      | M      | U         |
| F   | 82  | ET        | Nil                  | No                   | yes                 | U       | U      | U      | U         |
| M   | 78  | ET        | Nil                  | No                   | yes                 | U       | U      | U      | U         |
| F   | 79  | ET        | Nil                  | No                   | yes                 | U       | U      | U      | U         |
| M   | 84  | ET        | AML                  | No                   | yes                 | U       | U      | U      | U         |
| F   | 74  | ET        | Nil                  | No                   | yes                 | M       | U      | M      | U         |
| F   | 56  | ET        | Nil                  | No                   | yes                 | U       | U      | U      | U         |
| M   | 60  | ET        | Nil                  | No                   | N/A                 | U       | U      | U      | U         |
| M   | 68  | ET        | Nil                  | No                   | AML                 | U       | U      | U      | U         |
| M   | 39  | ET        | Nil                  | No                   | yes                 | U       | U      | U      | U         |
| F   | 79  | ET        | Weight loss          | No                   | yes                 | U       | U      | U      | U         |
| F   | 60  | ET        | erythromelalgia      | No                   | no                  | U       | U      | U      | U         |
| M   | 63  | ET        | Minor stroke (LUL numbness) | No | no | U | U | U | U |
| M   | 47  | ET        | erythromelalgia      | No                   | no                  | U       | U      | U      | U         |
| F   | 62  | ET        | erythromelalgia, Headache | No | no | U | U | U | U |
| F   | 83  | ET        | N/A                  | N/A (No record)      | No                  | U       | U      | M      | U         |
| F   | 32  | ET        | Nil                  | N/A (No record)      | no                  | U       | U      | U      | U         |
| F   | 42  | ET        | Nil                  | No                   | no                  | M       | U      | U      | U         |
| M   | 85  | ET        | Nil                  | No                   | no                  | U       | U      | U      | U         |
| M   | 71  | ET        | Nil                  | No                   | no                  | U       | U      | U      | U         |
| F   | 28  | ET        | Nil                  | No                   | no                  | U       | U      | U      | U         |
| F   | 87  | ET        | Nil                  | No                   | no                  | U       | U      | U      | U         |
| M   | 48  | ET        | Nil                  | No                   | no                  | U       | U      | U      | U         |
| M   | 48  | ET        | Nil                  | No                   | no                  | U       | U      | U      | U         |
| M   | 41  | ET        | epistaxis            | No                   | N/A                 | U       | U      | U      | U         |
| M   | 82  | ET        | N/A                  | No                   | N/A                 | U       | U      | U      | U         |
| F   | 41  | PMF       | Easy bruising        | N/A (No record)      | Yes                 | U       | U      | U      | U         |
| M   | 67  | PMF       | Nil                  | No                   | Yes                 | U       | U      | U      | U         |
| F   | 82  | PMF       | Nil                  | No                   | Yes                 | U       | U      | U      | U         |
|   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|
|   |   |   |   |   |   |   |   |
| F | 42 | PV | Minor stroke (RUL paraesthesia) | No | yes | U | U |
| M | 48 | PV | Gum bleeding | No | yes | U | U |
| F | 75 | PV | Nil | AML | yes | U | U |
| M | 78 | PV | Nil | No | yes | U | U |
| M | 57 | PV | Visual blurring (BRAO) | No | yes | U | U |
| M | 44 | PV | abdominal pain (splenic infarction) | No | no | U | U |
| M | 71 | PV | Gum bleeding | MDS | N/A | U | U |
| F | 77 | PV | N/A | No | N/A | U | U |

Abbreviations: ET: essential thrombocythemia; PV: polycythemia vera; PMF: primary myelofibrosis; CVA: cerebrovascular accident; BRAO: branch retinal vein occlusion; LUL: left upper limb; RUL: right upper limb; AML: acute myeloid leukemia; MDS: myelodysplastic syndrome; N/A: not available; U: unmethylated; M: methylated

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Table 2 MSP primer sequences and reaction conditions

| Gene   | Forward primer (5’ - 3’)                   | Reverse primer (5’ - 3’)                   | MgCl₂/Tm/Cycles | Reference |
|--------|--------------------------------------------|--------------------------------------------|-----------------|-----------|
| miR-34a |                                             |                                             |                 |           |
| M-MSP  | GGGGATGAGGATTAGGTTTTC                  | ACAAACGCAATAAAAAACGACG                  | 1.5 mM/58°C/35  | [9]       |
| U-MSP  | GGGGATGAGGATTAGGTTTTC                  | CAAACACACACATAAAAAACACAC                 | 1.5 mM/58°C/35  |           |
| miR-34b/c |                                            |                                            |                 |           |
| M-MSP  | ATTCTGGTCTTCTGGTTTTC                  | CGACTACAAGTGTTGTTTTTCTCTCTCA           | 2.0 mM/58°C/35  | [34,35]  |
| U-MSP  | TTTTTATTTTTTTTTTTTTTGTGTTTTTTTTGTG   | CAACACAGTCCAAAGACACATCC                 | 1.25 mM/56°C/38 |           |
| miR-124-1 |                                           |                                            |                 |           |
| M-MSP  | AAAGAAGTTTTTTGGAAGCTC                | AATAAAAAACGACGCCTATA                    | 1.5 mM/55°C/35  | [36]      |
| U-MSP  | AATAAAAAAGTTTTTTGGAAGATGGT             | AAAAAAATAAAAAACACACATAC                 | 2.0 mM/55°C/35  |           |
| miR-203 |                                             |                                            |                 |           |
| M-MSP  | GAGTATTTTACGCTTACGACGAGAC            | CTTTCTAGACGACAGCAGCC                   | 1.5 mM/58°C/35  | [37]      |
| U-MSP  | TTTGATGATTTTTGTTTATGAGATGAT           | AACACCTTTATACACACACACACCA               | 1.5 mM/58°C/35  |           |

Abbreviations: Tm, annealing temperature; M-MSP, MSP for the methylated allele; U-MSP, MSP for the unmethylated allele.

Statistical analysis

Correlation between combined miR methylation status (i.e. those with methylation of any of these four miRs) with categorical variables (gender, MPN subtype, occurrence of myeloid transformation, development of thrombosis [either at diagnosis or after diagnosis of MPN] and presence of JAK2 V617F mutation) and continuous variables (age, presenting Hb, leukocyte and platelet counts) was computed by the Chi-square test (or Fisher Exact test) and Student’s T-test. All p-values were two-sided.

Results

MSP

Controls

Direct sequencing of the M-MSP products from the methylated positive control confirmed the MSP specificity and complete bisulfite conversion, which methylated cytosine remained as cytosine upon sequencing (underlined) while unmethylated cytosine appeared as thymidine (Figure 1A). The positive and negative controls showed expected MSP results with normal DNA showing positive U-MSP but negative M-MSP amplification; and conversely, methylated control DNA showing negative U-MSP but positive M-MSP amplification. None of the 8 normal control marrows showed aberrant methylation of miR-34a, -34b/c, -124-1 or -203 (Figure 1B).

Cell lines

MSP analysis of the four cell lines showed that miR-34a was hemizygozously methylated in MEG-01 and K-562 and completely unmethylated in HEL and SET-2. miR-34b/c was completely methylated in HEL, hemizygously methylated in MEG-01 and completely unmethylated in K-562 and SET-2. miR-124-1 was completely unmethylated in all the four cell lines. miR-203 was completely unmethylated in K-562 and SET-2. However, there was neither U- or M-MSP signals of miR-203 in both HEL and MEG-01, suggesting a possibility of homozygous deletion (Table 3; Figure 1C).

Primary samples

In the 45 primary bone marrow samples, miR-34a was methylated in one (2.2%), miR-34b/c in four (8.9%), miR-203 in four (8.9%) of patients but none had methylation of miR-124-1 (Figure 2; Table 1). Moreover, two (4.4%) had concomitant methylation of miR-34b/c and -203 but none had concomitant methylation of miR-34a and -34b/c.

Direct sequencing of the methylated MSP products confirmed methylation of miRs in the primary samples (Figure 2). With regards to the MPN subtype, miR-34a methylation occurred in a patient with ET (2.9%), and miR-203 in four patients (11.8%) with ET. On the other hand, miR-34b/c methylation occurred in three patients (8.8%) with ET and one patient (12.5%) with PV. Overall, seven patients (15.6%) had methylation of at least one of the three miRs. miR methylation was not associated with age (p = 0.651), gender (p = 0.225), MPN subtype (p = 0.484), presenting Hb (p = 0.874), presenting leukocyte count (p = 0.969), presenting platelet count (p = 0.328), myeloid
A) Sequence analysis of the M-MSP product from bisulfite-treated positive control DNA showed that the cytosine (C) residues of CpG dinucleotides were methylated and remained unchanged, whereas all the other C residues were unmethylated and were converted to thymidine (T), confirming complete bisulfite conversion and MSP specificity. B) M-/U-MSP analysis showed that all the eight normal controls (N1-N8) were unmethylated. C) In the cell lines, MEG-01 and K-562 were hemizygously methylated for miR-34a; HEL was completely methylated, MEG-01 was hemizygously methylated for miR-34b/c; all the four cell lines were unmethylated for miR-124-1; K-562 and SET-2 were completely unmethylated for miR-203.

**Figure 1** Methylation of miR-34a, miR-34b/c, miR-124-1 and miR-203. A) Sequence analysis of the M-MSP product from bisulfite-treated positive control DNA showed that the cytosine (C) residues of CpG dinucleotides were methylated and remained unchanged, whereas all the other C residues were unmethylated and were converted to thymidine (T), confirming complete bisulfite conversion and MSP specificity. B) M-/U-MSP analysis showed that all the eight normal controls (N1-N8) were unmethylated. C) In the cell lines, MEG-01 and K-562 were hemizygously methylated for miR-34a; HEL was completely methylated, MEG-01 was hemizygously methylated for miR-34b/c; all the four cell lines were unmethylated for miR-124-1; K-562 and SET-2 were completely unmethylated for miR-203.
transformation (p = 0.99), thrombotic events (p = 0.311) or JAK2 V617F mutation (p = 0.99).

5-AzadC treatment of HEL cells

Untreated HEL cells were homozygously methylated for miR-34b/c (Chr. 11q23). After 5-AzadC hypomethylation treatment, miR-34b/c U-MSP signal emerged, together with about 3-fold increase in expression of both mature miR-34b and miR-34c on day 7 as analyzed by stem-loop RT-PCR (Figure 3). On the other hand, miR-34a was unmethylated in HEL. By conventional RT-PCR of the primary transcript of miR-34a, pri-miR-34a was not constitutively expressed, and treatment with 5-AzadC did not lead to expression of primary miR-34a (Additional file 1).

Discussion

Little information is available about the role of miRs in MPNs. Based on a literature search on the PubMed using keywords of "microRNA, methylation and myeloproliferative", no publication could be found. Therefore, this is likely the first report on methylation of miRs in MPN. In this study, we examined the methylation of miR-34a, -34b/c, -124-1 and -203 for a range of MPNs. In an attempt to identify miRs specifically involved in erythropoiesis, miR expression from in vitro expansion of erythroid cells derived from peripheral blood mononuclear cells were compared with controls, which showed miR-451 upregulation was specific to erythropoiesis [16]. On the other hand, during in vitro differentiation of megakaryocytes derived from CD34+ hematopoietic progenitors, downregulation of miR-10a, -10b, -17, -20, -106 and -126 was observed [17]. Moreover, when comparing the miR expression profiling of megakaryoblastic cell line with in vitro differentiated megakaryocytes, upregulation of miR-99a, -101, -126, and -135 was also found [17]. Therefore, the upregulation and down-regulation of these specific miRs were associated with megakaryocytic and erythropoietic differentiation.

Despite that multiple TSGs were shown to be silenced by hypermethylation in AML [4,14,18-22], methylation of tumor suppressor genes was less frequently reported in MPN. Only recently, there were a few reports of methylation of the negative regulators of the JAK-STAT signaling pathway, SOCS1, SOCS2 and SOCS3, in Ph-ve MPN regardless of the JAK2 V617F mutation status [23-26]. Furthermore, based on previous work by us and others, CDK6 has been shown to be the target of multiple miRs including miR-34a, -34b/c and -124-1 [27,28]. Moreover, CREB is the target of miR-34b and -203 [10,29]. Finally, ABL is an additional target of miR-203 [12]. Consequently, epigenetic silencing of tumor suppressor miR-34a, miR-34b/c, miR-124-1 and miR-203 will confer proliferative advantage to the tumor cells [27-29]. In contrast to a previous report which showed miR-203 was methylated in Ph+ve but not Ph-ve MPN or leukemia, using MSP primers in the similar region [12], we demonstrated that miR-203 was hypermethylated in primary MPN samples, which was further verified by direct sequencing of the methylated MSP products. Therefore, it would appear that miR-203 methylation is involved in a wider spectrum of MPNs or leukemias, regardless of their Ph chromosome status. Finally, while two patients had concomitant methylation of miR-203 and -34b/c, none had concomitant methylation of miR-34a and -34b/c, both transcriptional targets of p53, and hence avoiding duplication of tumor suppressor gene inactivation of the same pathway.

In HEL and MEG-01 cells, both U- and M-MSP signals of miR-203 were absent, which might be due to the following possibilities: (1) sample DNA degradation, (2) inappropriate PCR condition, or (3) homozygous deletion of the region. Since simultaneous U-MSP analysis of the same DNA sample for miR-34a, miR-34b/c, and miR-124-1 promoter successfully generated the U-MSP signals, hence the absence of MSP signals for miR-203 in HEL and MEG-01 cells could not be explained by a poor DNA quality. Moreover, as miR-203 U-MSP was successful in all the other samples including cell lines, normal controls, patient samples and methylated positive control, inappropriate MSP conditions appears unlikely. Therefore, the absence of both M- and U-MSP signals in HEL and MEG-01 cells might be caused by deletions of the region. However, karyotypic data of HEL and MEG-01 cells did not reveal homozygous deletion of 14q32, and hence whether absence of MSP amplification of miR-203 might be due to microdeletion of this region requires further study [30]. Furthermore, hypomethylation treatment of the HEL cells, which was homozygously methylated for the miR-34b/c, a microRNA cluster localized to 11q23, showed significant re-expression of mature miR-34b and miR-34c. This

Table 3 MSP of miRs in cell lines

| Chromosomal location of miR | miR-34a | miR-34b/c | miR-124-1 | miR-203 |
|----------------------------|---------|-----------|-----------|---------|
| HEL                        | 1p36    | 11q23     | 8p23      | 14q32   |
| MEG-01                     | UM      | MM        | UU        | -/-     |
| K-562                      | UM      | UM        | UU        | -/-     |
| SET-2                      | UU      | UU        | UU        | UU      |

UU: completely unmethylated; UM: hemizygously methylated; MM: homozygously methylated; -/-: absence of both U- and M-MSP signals
Figure 2. Methylation of miR-34a, miR-34b/c, miR-124-1 and miR-203 in primary samples. For each of these miRs, M-/U-MSP and sequencing of the M-MSP product from representative bisulfite-treated primary samples were shown. In the sequence analysis of the M-MSP product, cytosine [C] residues of CpG dinucleotides were methylated and remained unchanged, partially methylated C residues were denoted as [C/T], unmethylated C residues were converted into [T], whereas all the non-CpG C residues were unmethylated and were converted to thymidine [T], confirming complete bisulfite conversion and MSP specificity.
finding is consistent with that both miR-34b and miR-34c are under the promoter regulation of the same CpG island. By contrast, miR-34a, another member of the miR-34 family localized to 1p36, was not constitutively expressed. Moreover, hypomethylating treatment did not lead to expression of primary miR-34a, suggesting additional mechanism, possibly histone modification, in the regulation of miR-34a expression [9]. Furthermore, in addition to showing miR silencing in cell line, it is important to show the correlation of miR methylation and miR expression in the primary sample. In this connection, we shall collect both DNA and RNA from diagnostic bone marrow samples in the future.

In order to identify miRs that are methylated specifically at the time of transformation to AML or MDS, and hence implicated for pathogenesis of myeloid transformation, ideally one should analyze the paired marrow samples at the time of transformation to AML or MDS, and hence implicated for pathogenesis of myeloid transformation.

Finally, unlike the association of TSG with clinical parameters, such as the association of CDKN2B and WIF1 methylation with high presenting leukocyte count in acute promyelocytic leukemia [32,33], methylation of these miRs did not correlate with demographic, presenting blood counts, JAK2 V617F mutation or complications including thrombosis and myeloid transformations.

**Conclusion**

This is the first report of miR hypermethylation in MPNs. miR-203 hypermethylation is not specific to Ph+ve leukemias but also present in Ph-ve MPNs. miR-34b/c methylation was associated with reversible miR silencing. There was no correlation of miR methylation with clinical demographic data or outcome.

**Additional material**

Additional file 1: Effect of 5-aza-2’-deoxycytidine (5-AzadC) treatment on HEL cells. 5-AzadC treatment of HEL cells.

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**Authors’ contributions**

All authors read and approved the final manuscript. CSC is responsible for design of study, interpretation of data and writing manuscript. TSW is responsible for provision of study material. TFW is responsible for provision of cell lines. KFW is responsible for provision of patients’ material.

**Competing interests**

The authors declare that they have no competing interests.

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