Efficacy of Azacitidine in Autoimmune and Inflammatory Disorders associated with Myelodysplastic Syndromes and Chronic Myelomonocytic Leukemia

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Efficacy of Azacitidine in Autoimmune and Inflammatory Disorders associated with Myelodysplastic Syndromes and Chronic Myelomonocytic Leukemia

Running title: Azacitidine in autoimmune disorders associated with MDS/CMML

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Highlights

1) AID associated to MDS/CMML is difficult to manage

2) Azacitidine can improve the control of AID associated with MDS/CMML.

3) Azacitidine allows to reduce steroid dose in most MDS patients with concurrent AID

4) Azacitidine can be beneficial on AID, even when inactive on the underlying MDS/CMML.

Summary

This retrospective study describes efficacy of Azacitidine on autoimmune disorders (AID) associated with MDS/CMML in 22 patients. Response of AID to Azacitidine was observed in 19 patients (86%). Reduction or discontinuation of steroids and/or immunosuppressive therapy (IST) was possible in 16 cases (73%). Hematologic response was seen in 55% of the patients. MDS/CMML and AID evolution was concordant in 13 cases (59%): both favorable (n=11), both unfavorable (n=2), but AID improved while MDS/CMML worsened (n=8) and vice versa (n=1). Azacitidine frequently seems effective in controlling steroid-dependent AID associated with MDS/CMML, but prospective studies are necessary to confirm those findings.

Key words: Azacitidine, myelodysplastic syndromes, autoimmune disorders.
**Introduction**

Autoimmune disorders are observed in 10 to 30% of MDS and CMML, typically diagnosed concomitantly or shortly before or after MDS/CMML [1].

Most common AID associated with MDS/CMML include relapsing polychondritis, vasculitis, non-erosive and seronegative arthritis [2,3] and Sweet’s syndrome [4]. While AID associated with MDS/CMML usually respond to corticosteroids [2], many patients remain steroid dependent or resistant, requiring additional IST, a situation which increases the risk of severe cytopenias and infections in the context of MDS/CMML [1].

Azacitidine significantly improves survival in higher risk MDS [5] and in myelodysplastic type CMML [6,7] and is also approved outside Europe for the treatment of lower risk MDS. Seven patients with MDS/CMML and AID treated with Azacitidine have been reported to our knowledge [8–11].

**Methods**

We retrospectively analyzed the efficacy of Azacitidine in MDS/CMML patients with concomitant AID seen between 2007 and May 2014 in registries of the Société Nationale Française de Médecine Interne (SNFMI) and of the Groupe Francophone des Myélodysplasies. The SNFMI registry included 123 MDS/CMML patients with AID in 26 centres) [13], while the GFM registry included 4202 MDS/CMML in 30 centres of whom 80 also had documented AID. The GFM registry has been so far the basis for many publications of the GFM or that included the GFM [14–16].

Inclusion criteria for the present study were (i) complete or atypical AID according to usual international criteria [17], (ii) MDS or non-proliferative CMML (ie with WBC<13G/L), classified according to WHO 2008 criteria [18] but also including
patients with up to 30% marrow blasts, (iii) treatment with at least one cycle of Azacitidine.

Patients were excluded if AID were caused by infectious disease, AID were in remission without steroids or IST at Azacitidine onset, or if IST was administered more than 12 months before diagnosis of MDS, suggesting MDS secondary to IST.

Complete/ partial remission (CR/PR) of AID after Azacitidine were defined by complete/partial disappearance of clinical or biological signs with stable dose or decrease/discontinuation of steroids or IST, as previously defined [2]. Hematological response to Azacitidine was classified according to IWG 2006 criteria [18].

For statistical analysis, Wilcoxon test was used for analysis of paired quantitative variables and Fisher's exact test for paired qualitative variables using R 3.2.1 (2015) with package Rcmdr 2.1-7 (2007).

Results

Among patients with both MDS/CMML and AID included in the 2 registries, twenty eight patients had received Azacitidine (flow chart in figure 1). Six were excluded, because AID were in remission without IST at Azacitidine onset (n=2), of inadequate follow-up (n=3) and as vasculitis was post-infectious (n=1). Median age of the 22 remaining patients was 70 years (range 41-84), including 6 females and 16 males (73%). (Table 1)

Diagnosis of MDS/CMML preceded AID diagnosis (n=8; by a median of 17 months), was concomitant with (n=7) or followed AID diagnosis (n=7; by a median of 20 months).
**AID and MDS characteristics**

At MDS/CMML diagnosis, 14 patients had low or int-1 IPSS and 8 had int-2 or high IPSS. The 2 cases of CMML with WBC below 13G/L were classifiable by IPSS. AID diagnosis included: Behçet’s disease (n=4), polymyalgia rheumatica (n=3), polymyalgia rheumatica with giant cell arteritis (n=1), giant cell arteritis (n=1), relapsing polychondritis (n=3), polychondritis with Sweet’s syndrome (n=1), Sweet’s syndrome (n=1), systemic lupus erythematosus (n=2), seronegative polyarthritis (n=2), Sjögren’s syndrome with anti-phospholipid syndrome (n=1), adult onset Still’s disease (n=1), large vessel vasculitis (n=1) and unclassified small vessel vasculitis (n=1). Twelve (55%) AID were considered atypical and 10 (45%) fulfilled complete diagnostic criteria.

**AID features at Azacitidine onset**

At Azacitidine onset, AID was still active (n=15), in PR (n=5) and in CR (n=2). Nineteen patients still reported clinical symptoms related to AID including asthenia (n=17), weight loss (n=8), fever >38.5°C (n=8), rheumatologic signs (n=15), skin involvement (n=10), oral aphtosis (n=4) (Table 1). Twenty (91%) patients were receiving steroids (median: 23 mg/day) for a median duration of 11 months (range 1-101) with no efficacy (n=13), partial efficacy (n=5) and full efficacy on AID (n=2). In patients where steroids were ineffective, they were however generally continued to avoid AID flare. The median duration of steroid use, in the 18 good responders on AID, was 16 months (range 1-101) versus 9 (range 8-9) in the non-responders. Thirteen patients were non-responders to IST and 11 received concomitant treatment with IST and steroids: without efficacy in 7 patients, partial efficacy in 2 and full efficacy in 2 (Table 1). Onset of IST preceded that of Azacitidine by a median of 7
months (range 1-50); the 3 patients who did not have a good response on AID did not receive IST.

Median interval between MDS/CMML diagnosis and Azacitidine onset was 9 months (range 1-93). Azacitidine was the first-line treatment of MDS/CMML in 16 patients (73%) and was in 6 patients preceded by lenalidomide (n=2), hydroxyurea (n=2), idarubicin (n=1) and low dose cytarabine (n=1), treatments to which AID had not responded in any patient.

**MDS features at Azacitidine onset**

At Azacitidine onset, 11 patients had int-2 or high IPSS, and Azacitidine was used according to the drug label in EU; in the 11 low or int-1 IPSS, reasons to start Azacitidine were RBC transfusion dependence or severe thrombocytopenia in 8 patients (6 of whom also had AID flare) and AID flare alone in 3 patients (Table1).

**AID response**

Patients received a median of 6 cycles of Azacitidine (range 3-14). CR of AID was observed in 16 patients (73%), PR in 3 cases (14%) and no effect or worsening of AID in 3 cases (14%). All responses were observed within 3 cycles, but 8 PR after 3 cycles became CR after 6 months. In the 3 patients treated with Azacitidine for AID flare alone, one achieved CR, one PR and the last patient did not respond. Steroids were tapered from 40 mg/day to 16 mg/day in one patient and IST was discontinued in another patient.

Median prednisone dose was reduced from 23 mg/day to 9 mg/day (p=0.001), including prednisone discontinuation in 3 patients (14%) and dose reduction in 14 patients (64%) (Table1). IST discontinuation was possible in 7 patients (32%). In the 9 steroid-dependent or resistant patients, median prednisone dose was decreased from 50mg/day to 5mg/day (p=0.01) and prednisone discontinuation was possible in
3 patients. IST discontinuation was possible in the 3 patients who received them. After Azacitidine treatment, median CRP levels decreased from 40 (range 0-300) to 15 mg/L (range 0-146) (p=0.28).

**MDS response**

Response of MDS/CMML to Azacitidine was seen in 12 patients (55%), including CR in 9 patients (41%), PR in 1, stable disease with erythroid response in 1, marrow CR in 1, stable disease without response of cytopenias in 8 and progression in 2.

MDS/CMML and AID evolution were concordant in 13 cases (59%): both improved (n=11) or worsened (n=2). In the other patients, AID improved while MDS worsened (n=8) and vice versa (n=1).

**Side effects/ Evolution**

Adverse events occurring during Azacitidine treatment included: 9 grade 2-3 infections (6 of them occurred in Int-2 or high IPSS patients), 1 fatal CNS bleeding, 1 deep vein thrombosis, 1 Interstitial lung disease.

Only 3 relapses of AID were seen after 3, 19 and 19 months respectively, and 5 patients had response duration ≥ 12 months, but median follow-up from response to Azacitidine was only 8 months (Table1).

Seven patients (32%) progressed to AML and 10 patients (45%; including 80% of IPSS int-2 and high risk patients) had died, from CNS bleeding (n=3), severe infection (n=5), progression in AML (n=1) and myocardial infarction (n=1). Among the 10 patients who died, 6 had progressed to AML. Severe infections were more frequently linked to AML progression than to Azacitidine. Median survival from Azacitidine onset was 16 months in IPSS high or int-2 MDS, and was not reached in IPSS low and int-1 MDS.
Discussion

Azacitidine improved clinical signs of AID in 86% of the cases (73%CR, 14%PR) in patients with MDS/CMML. Azacitidine also allowed to reduce the steroid dose in 64% of the patients and discontinuation of other IST in 32% of them.

In patient n°5, Azacitidine allowed to discontinue both MMF and hydroxychloroquine without having an AID flare. In patient n°10, prolonged clinical and biological complete response was maintained under Azacitidine without requirement of new IST or steroids.

In the only 7 cases of AID with MDS/CMML treated with Azacitidine published so far to our knowledge, 6 were refractory to steroids and/or IST and 4 had higher risk IPSS. MDS and AID evolution were concordant in six cases (Table 2) [8–11].

In our series, all responses of AID to Azacitidine were seen by the third cycle, although about one third of responses improved and became complete between 3 and 6 cycles. Furthermore, although less than 50% of the patients received more than 6 cycles of Azacitidine, only 3 relapses were observed (but with a median follow-up from response of AID to Azacitidine of only 8 months).

Most of the MDS/CMML included in the present study required some treatment because of high or int-2 IPSS in 11 cases, RBC transfusion dependence or severe thrombocytopenia in 8 cases. In the last 3 patients, however AID flare refractory to IST and/or high dose steroids was the only reason to start Azacitidine.

The concordant evolution of AID and MDS/CMML we observed in 59% of the cases further supports a pathophysiological link between MDS/CMML and AID. However, it often remains unclear, in MDS/CMML associated with AID, which of the 2 diseases is the cause and which is the consequence. While immunological defects may potentially trigger MDS/CMML, they may also be in some cases the consequences of
MDS/CML, when some cells of the immune system are part of the MDS/CML clone [1,11]. In addition to its direct pro-apoptotic activity on MDS cells, Azacitidine also has immunomodulatory functions which could explain at least in part its effect in MDS-related AID. Azacitidine can indeed expand CD4+CD25+/FOXP3+ regulatory T cells which are decreased with impaired suppressive function in many AID [19]. Azacitidine also increases expression of anti-inflammatory molecules like IFNγ and could inhibit CD4+ T-cells and reduce immune-mediated cytotoxicity [11]. Azacitidine could finally decrease the synthesis of inflammatory cytokines like IL-6 [20]. In some cases, Azacitidine therefore could have a positive immunomodulatory effect, even when it is not active on MDS/CML.

In conclusion, our results suggest that Azacitidine can improve the control of AID associated to MDS/CML and allow to reduce steroid dose in most patients with MDS and concurrent AID. Prospective studies are necessary to confirm this beneficial effect and the safety of Azacitidine in AID associated with MDS/CML.

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Contributions

FJB performed the research; MA, FP, FO and BT designed the research study; FJB analysed the data; GE, KJE, AJB, DO, DG, BAL, OM, MG, AA, LN, BS, LE, PS, GC, LO, RJ, FP, FO and BT share data on patients; and FJB wrote the paper.

ACKNOWLEDGMENT

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References


FIGURE 1

GFM registry (n=4202)
- MDS/CMML without AID (n=4122)
  - MDS/CMML with AID (n=80)
    - Untreated with AZA (n=67)
  - MDS/CMML with AID treated by AZA (n=13)
    - Inadequate follow up (n=3)*
    - AID remission at AZA onset (n=1)*
  - Included (n=9)

SNFMI registry (n=123)
- MDS/CMML with AID
  - Untreated with AZA (n=108)
  - MDS/CMML with AID treated by AZA (n=15)
    - Included in GFM (n=3)
    - Inadequate follow up (n=2)*
    - AID remission at AZA onset (n=2)*
  - Included (n=8)

Patients included (n=22)
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<th>IST Before/after AZA</th>
<th>CRP (mg/L) before/after AZA</th>
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<td>RAEB1 46,XY</td>
<td>Low 59</td>
<td>PC</td>
<td>IFX ANA</td>
<td>F</td>
<td>As Sk Ji</td>
<td>25/30</td>
<td>ANA/ None</td>
<td>50/60</td>
<td>CR</td>
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<td>16</td>
<td>M</td>
<td>57</td>
<td>RA 46,XY</td>
<td>Low 26</td>
<td>PC</td>
<td>IFX MTX CYC</td>
<td>F</td>
<td>As Sk Ji ENT</td>
<td>15/18</td>
<td>None/ None</td>
<td>248/10</td>
<td>CR</td>
<td>PR</td>
<td>CNS 9+</td>
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<tr>
<td>17</td>
<td>M</td>
<td>67</td>
<td>Del-5q MDS 46,XY,del(5q)</td>
<td>Int-1 93</td>
<td>PC &amp; Sweet</td>
<td>CIC</td>
<td>F</td>
<td>WL As Fe Sk Ji Lu ENT</td>
<td>80/5</td>
<td>None/ None</td>
<td>240/NA</td>
<td>CR</td>
<td>CR</td>
<td>None 8+</td>
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</tr>
<tr>
<td>18</td>
<td>M</td>
<td>66</td>
<td>RAEB1 46,XY</td>
<td>Int-1 0</td>
<td>GCA</td>
<td>ANA IFX MTX TCZ IVIG CIC</td>
<td>F</td>
<td>As Sk ENT</td>
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<td>IVIG/ None</td>
<td>146/22</td>
<td>CR</td>
<td>CR</td>
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<td>19</td>
<td>M</td>
<td>84</td>
<td>RCMD 46,XY</td>
<td>Int-1 8</td>
<td>PMR</td>
<td>Col</td>
<td>F</td>
<td>As Sk Ji</td>
<td>17/15</td>
<td>None/ None</td>
<td>61/143</td>
<td>mCR</td>
<td>NR</td>
<td>Fe Ji 3</td>
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<tr>
<td>20</td>
<td>M</td>
<td>47</td>
<td>RCMD 47,XY, +8</td>
<td>Int-1 9</td>
<td>Still</td>
<td>Col MTX ANA TCZ HCO</td>
<td>F</td>
<td>WL As Fe, Ji ENT PN</td>
<td>40/16</td>
<td>ANA, HCO/ None</td>
<td>38/NA</td>
<td>CR</td>
<td>PR</td>
<td>Ji 2+</td>
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<tr>
<td>21</td>
<td>M</td>
<td>64</td>
<td>RAEBt 46,XY</td>
<td>Int-2 2</td>
<td>PC</td>
<td>ETC</td>
<td>F</td>
<td>Ji</td>
<td>80/5</td>
<td>None/ None</td>
<td>161/14</td>
<td>Prog</td>
<td>CR</td>
<td>None 31+</td>
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<tr>
<td>22</td>
<td>M</td>
<td>83</td>
<td>RA 46,XY</td>
<td>Int-1 11</td>
<td>PC</td>
<td>AZA</td>
<td>F</td>
<td>Ji</td>
<td>14/10</td>
<td>AZAT/None</td>
<td>225/5</td>
<td>CR</td>
<td>CR</td>
<td>None 3+</td>
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Table 2: Published cases of MDS/CMML with AID treated with Azacitidine

<table>
<thead>
<tr>
<th>Gender, Age</th>
<th>MDS/CMML</th>
<th>IPSS</th>
<th>AID</th>
<th>IST failure</th>
<th>Evolution</th>
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<tbody>
<tr>
<td>1, M 74[8]</td>
<td>RAEB2</td>
<td>High</td>
<td>Sweet</td>
<td>CST, HCQ</td>
<td>CR of MDS and AID</td>
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<tr>
<td>2, M 44[9]</td>
<td>RCMD</td>
<td>Int-2</td>
<td>SLE</td>
<td>CST(40/d), colchicine, disulone, thalidomide</td>
<td>CR of MDS and AID</td>
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<tr>
<td>3, M 69[10]</td>
<td>RCMD</td>
<td>Int-1</td>
<td>Polyarthritis &amp; Sweet</td>
<td>CST(8mg/d)</td>
<td>CR of MDS and AID CST(8mg/d)</td>
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<tr>
<td>4, M, 66[10]</td>
<td>CMML</td>
<td>Int-1</td>
<td>SS &amp; APLS</td>
<td>CST(20mg/d), MMF, MTX, CYC,</td>
<td>CR of MDS and AID CST(5mg/d) MMF stopped</td>
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<tr>
<td>5, M, 64[10]</td>
<td>RAEB-1</td>
<td>Int-1</td>
<td>Polyarthritis &amp; Sweet</td>
<td>CST(10mg/d), thalidomide.</td>
<td>CR of MDS and AID CST stopped</td>
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</tbody>
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