Accumulating evidence implicates innate immune activation in the pathobiology of myelodysplastic syndromes. A key myeloid-related inflammatory protein, S100A9, serves as a Toll-like receptor ligand regulating tumor necrosis factor-α and interleukin-1β production. The role of myelodysplastic syndrome-related inflammatory proteins in endogenous erythropoietin regulation and response to erythroid-stimulating agents or lenalidomide has not been investigated. The HepG2 hepatoma cell line was used to investigate in vitro erythropoietin elaboration. Serum samples collected from 311 patients with myelodysplastic syndrome were investigated (125 prior to treatment with erythroid-stimulating agents and 186 prior to lenalidomide therapy). Serum concentrations of S100A9, S100A8, tumor necrosis factor-α, interleukin-1β and erythropoietin were analyzed by enzyme-linked immunosorbent assay. Using erythropoietin-producing HepG2 cells, we show that S100A9, tumor necrosis factor-α and interleukin-1β suppress transcription and cellular elaboration of erythropoietin. Pre-incubation with lenalidomide significantly diminished suppression of erythropoietin production by S100A9 or tumor necrosis factor-α. Moreover, in peripheral blood mononuclear cells from patients with myelodysplastic syndromes, lenalidomide significantly reduced steady-state S100A9 generation (P=0.01) and lipopolysaccharide-induced tumor necrosis factor-α elaboration (P=0.002). Enzyme-linked immunosorbent assays of serum from 316 patients with non-del(5q) myelodysplastic syndromes demonstrated a significant inverse correlation between tumor necrosis factor-α and erythropoietin concentrations (P=0.006), and between S100A9 and erythropoietin (P=0.01). Moreover, baseline serum tumor necrosis factor-α concentration was significantly higher in responders to erythroid-stimulating agents (P=0.03), whereas lenalidomide responders had significantly lower tumor necrosis factor-α and higher S100A9 serum concentrations (P=0.05). These findings suggest that S100A9 and its nuclear factor-kB transcriptional target, tumor necrosis factor-α, directly suppress erythropoietin elaboration in myelodysplastic syndromes. These cytokines may serve as rational biomarkers of response to lenalidomide and erythroid-stimulating agent treatments. Therapeutic strategies that either neutralize or suppress S100A9 may improve erythropoiesis in patients with myelodysplastic syndromes.
Introduction

Ineffective erythropoiesis in patients with myelodysplastic syndromes (MDS) derives from both intrinsic abnormalities affecting the response to erythropoietin and extrinsic pressures on the inflammatory bone marrow microenvironment. A number of inflammatory cytokines such as tumor necrosis factor-α (TNFα), interleukin (IL)-1β, IL-6 and others, are generated in excess in MDS and adversely influence hematopoietic stem and progenitor cell survival. Moreover, in a subset of MDS patients, endogenous erythropoietin production is deficient, further compromising erythropoietic potential. Accumulating evidence implicates innate immune activation in the pathophysiology of MDS and the accompanying inflammatory microenvironment. Bone marrow plasma concentrations of the pro-inflammatory, danger-associated molecular pattern (DAMP) protein, S100A9, are profoundly increased in lower-risk MDS, which serves as a catalyst directing myeloid-derived suppressor cell expansion. S100A9 is a ligand for CD33 and the Toll-like receptor (TLR)-4 which, through nuclear factor-kB (NF-kB) activation, regulates the transcription and cellular elaboration of inflammatory cytokines such as TNFα and IL-1β. The latter cytokines have been shown to suppress erythropoietin elaboration and have been implicated in the suppression of endogenous erythropoietin production in patients with anemia of chronic inflammation. The involvement of 100A8/S100A9 in del(5q) MDS has already been described. Erythropoietin-stimulating agents (ESA) and lenalidomide are efficient treatments used in lower-risk myelodysplastic syndromes. To date, the role of inflammatory parameters in the regulation of endogenous erythropoietin production and response to erythropoietic treatments in MDS has not been investigated. Here we show the importance of these inflammatory cytokines as key biological determinants of endogenous erythropoietin production and response to ESA and lenalidomide treatments in patients with non-del (5q) MDS.

Methods

Reagents and antibodies

Recombinant S100A9 was generated as previously described. TNFα, IL-1β and lipopolysaccharide were purchased from Sigma-Aldrich (Saint Louis, MO, USA). Lenalidomide was purchased from Fisher Scientific (Pittsburgh, PA, USA). A CD3 chimera was constructed as described elsewhere. NF-kB and Rho GDI antibodies were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Lamin A/C was purchased from Cell signaling (Saint Quentin, France). BMS444541 was purchased from Tocris Bioscience (Bristol, UK).

Cell culture

HepG2 cells, acquired from the American Type Culture Collection (ATCC, Manassas, VA, USA), were grown in Eagle’s minimum essential medium supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin solution. Cells were maintained at 37°C under 5% CO2.

Enzyme-linked immunosorbent assays

Human S100A9/MRP14 in patients’ serum and supernatants of the HepG2 cell line was quantified using a CircuLex S100A9/MRP14 enzyme-linked immunosorbent assay (ELISA) Kit (MBL, Nagano, Japan). Quantitative measurements of human TNFα and IL-1β were made using the Human TNFα ELISA Kit and Human IL-1β ELISA Kit, respectively (Life Technologies, Carlsbad, CA, USA). Human erythropoietin in patients’ serum and HepG2 cell line supernatants was quantified using a Human Erythropoietin ELISA Kit (Stemcell Technologies, Vancouver, BC, Canada). All measurements were performed in duplicate.

Real-time quantitative polymerase chain reaction

RNA was isolated using the RNeasy Mini Kit (Qiagen, Valencia, CA, USA) followed by iScript cDNA synthesis (Bio-Rad, Hercules, CA, USA) and amplification using iQ SYBR Green Supermix (Bio-Rad, Hercules, CA, USA). The relative level of gene expression for each experimental sample was calculated using the ΔΔCt method. Untreated cells were the experimental control and the housekeeping gene GAPDH was the endogenous control.

Western blot analysis

After treatment for 24 h, cells were harvested and lysed in 1X RIPA buffer supplemented with protease and phosphatase inhibitors for classical western blotting. For the nuclear extraction, cells were lysed in ice with buffer A, then pelleted. After removal of supernatant (cytoplasmic fraction), pellets were lysed in ice with buffer B and pelleted (nuclear fraction) (Nuclear Extraction Kit, Abcam, Cambridge, USA). Lysates were pelleted and 50 µg of protein were resolved by sodium dodecyl sulfate polyacrylamide gel electrophoresis and transferred to polyvinylidene fluoride membranes. The membranes were blocked for 30 min in 5% non-fat dry milk solution in PBST (phosphate-buffered saline with 0.1% Tween 20) and incubated with the indicated antibodies. Membranes were developed using ECL according to the manufacturer’s protocol (GE Healthcare, Little Chalfont, UK). Densitometry analysis was performed using Image J Software.

Patients and serum samples

Serum samples for ELISA analysis were collected from four centers (Taussig Cancer Institute, Cleveland, USA; AOU Careggi, University of Florence, Italy; Saint Louis Hospital, Paris, France; H. Lee Moffitt Cancer Center, Tampa, FL, USA). Peripheral blood mononuclear cells were collected from patients at Moffitt Cancer Center. All patients had provided consent to Institutional Review Board, or equivalent, approved protocols in hematology clinics at each center, and the Eastern Cooperative Oncology Group (ECOG) E2905 trial (www.clinicaltrial.gov NCT00843882). All routine clinical and biological data were available.

Statistical analysis

Data are expressed as mean ± standard error for continuous variables, or percentage of total for non-continuous variables. Spearman correlation, Mann-Whitney and Jonckheere-Terpstra tests were used for analysis of continuous variables. The chi-square test was used for analysis of non-continuous variables. Differences between the results of comparative tests were considered statistically significant if the two-sided P-value was less than 0.05. All statistical analyses were performed using SPSS v.22 software (IBM SPSS Statistics).

Results

Inflammatory proteins suppress erythropoietin production by HepG2 cells

Hepatoma HepG2 cells, which produce erythropoietin under basal conditions, were treated with varied concentrations of each of the inflammatory proteins, TNFα, IL-1β...
or S100A9. After 24 h exposure, we observed a concentration-dependent reduction in erythropoietin elaboration (Figure 1). At a concentration of 10 ng/mL, TNFα yielded a 40% reduction in erythropoietin elaboration (Figure 1A), compared to a 60% reduction following incubation with IL-1β (Figure 1B). Concentrations of 10 to 20 μg/mL of S100A9 completely suppressed erythropoietin elaboration, while 1 μg/mL yielded a 95% reduction (Figure 1C). For subsequent experiments, concentrations of 10 ng/mL of TNFα, 10 ng/mL of IL-1β and 1 μg/mL of S100A9 were employed.

Lenalidomide mitigates suppression of erythropoietin production by S100A9 and tumor necrosis factor-α

The transcription factor NF-κB is activated by TLR ligands and inflammatory cytokines such as TNFα which, like GATA2, is implicated in transcriptional suppression of the erythropoietin transcript. HepG2 cells are known to express the S100A9 receptor, TLR4, on the plasma membrane. Lenalidomide has been reported to suppress NF-κB activation in response to inflammatory cytokine stimulation in lymphocytes and other cell lineages. To determine whether lenalidomide can modulate suppression of erythropoietin production by inflammatory proteins, we treated HepG2 cells with 1 μM lenalidomide or vehicle control for 30 min prior to exposure to TNFα, IL-1β or S100A9. Lenalidomide significantly, but in incomplete-

Figure 1. Effects of tumor necrosis factor-α, interleukin-1β and S100A9 on erythropoietin elaboration in the HepG2 cell line. HepG2 cells were stimulated for 24 h with the indicated concentrations of (A) TNFα, (B) IL-1β and (C) S100A9 and erythropoietin (EPO) elaboration was determined by ELISA.
ly, reversed suppression of erythropoietin production by S100A9 (91% suppression by S100A9 versus 65% after lenalidomide pre-incubation, \( P=0.04 \)). Following treatment with TNF\( \alpha \), lenalidomide pre-treatment abrogated cytokine suppression of erythropoietin elaboration (TNF\( \alpha \), 67% suppression versus lenalidomide pre-incubation, 18%; \( P=0.05 \)). Lenalidomide had no effect on IL-1\( \beta \)-directed suppression of erythropoietin elaboration (Figure 2A). We also found that BMS-344541, a specific inhibitor of NF-\kappaB, abrogated cytokine suppression of erythropoietin elaboration with the same conditions (Figure 2A). Moreover, lenalidomide decreased basal TNF\( \alpha \) release after lenalidomide stimulation (Figure 2B).

IL-10 is known to be an anti-inflammatory cytokine secreted by immune cells.\(^{20}\) We performed quantitative polymerase chain reaction analysis to determine the effects of the inflammatory proteins and lenalidomide on IL-10 gene transcription. Pre-incubation of each inflammatory cytokine with lenalidomide significantly increased IL-10 mRNA expression compared to S100A9 or TNF\( \alpha \) treatment alone (Figure 2C). Lenalidomide had no modulatory effect on IL-10 following IL-1\( \beta \) treatment (data not shown).

Finally, NF-\kappaB is a key transcription factor involved in S100A9 and TNF\( \alpha \) receptor signaling and the transcriptional suppression of erythropoietin mRNA in erythropoietin-producing cells which is modulated by lenalidomide.\(^{21,22}\) We showed that NF-\kappaB targets, including TNF\( \alpha \) and IL-10, were regulated by lenalidomide. We therefore performed cytoplasmic and nuclear NF-\kappaB western blotting in HepG2 cells to discern the effect of lenalidomide on inflammatory protein signaling. As demonstrated by nuclear localization after lenalidomide pre-incubation, active NF-\kappaB was significantly reduced following treatment with S100A9 or TNF\( \alpha \), indicating that lenalidomide suppressed NF-\kappaB activation (Figure 2D). To confirm the results observed in HepG2 cells, we investigated the effects of lenalidomide on steady-state production of S100A9 by peripheral blood mononuclear cells isolated from patients with non-del(5q) MDS (n=7). ELISA showed a significant reduction in S100A9 elaboration after 24 h exposure to lenalidomide (\( P=0.01 \)) (Figure 3A). Similarly, pre-incubation of MDS peripheral blood mononuclear cells with lenalidomide significantly reduced TNF\( \alpha \) production induced by lipopolysaccharide (\( P=0.002 \)). These findings indicate that lenalidomide-modulated S100A9 and TNF\( \alpha \) suppression of erythropoietin elaboration is NF-\kappaB-dependent.

**Relationship between inflammatory proteins and endogenous erythropoietin concentration in patients with myelodysplastic syndromes**

To validate the regulatory role of inflammatory proteins on erythropoietin elaboration in vivo, we assessed the relationships between serum concentrations of various inflammatory cytokines and erythropoietin in MDS patients with symptomatic anemia. Serum samples from 316 patients with non-del(5q) MDS were analyzed. The median age of the patients was 74.7 years (range, 41-94). Distribution of International Prognostic Scoring System (IPSS) categories was low, intermediate-1, intermediate-2 and high risk in 38%, 50%, 10% and 2% of patients, respectively; whereas 24%, 38%, 22%, 13% and 3% of patients were very low, low, intermediate, high and very high risk according to the revised IPSS (IPSS-R) (Table 1).

Serum concentrations of erythropoietin, S100A9, S100A8, TNF\( \alpha \) and IL-1\( \beta \) were assessed by ELISA. The serum S100A9 concentration was significantly higher in patients with lower-risk MDS than in those with higher-risk MDS (12,226 pg/mL (range, 0-228,880) versus 240 pg/mL (range, 0-43,585), respectively; \( P=0.001 \). No significant differences were observed in TNF\( \alpha \) and IL-1\( \beta \) concentrations in IPSS-R category (data not shown).

There was a statistically significant negative correlation between TNF\( \alpha \) and erythropoietin concentrations (\( r = -0.240 \)).

**Table 1. Patients’ demographics and disease characteristics.**

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<th>Prior to ESA treatment (n=159)</th>
<th>Prior to lenalidomide ± ESA treatment (n=159)</th>
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**Figure 3. Effect of lenalidomide on S100A9 and tumor necrosis factor-\( \alpha \) production in peripheral blood mononuclear cells from patients with non-del(5q) myelodysplastic syndrome.** (A) Frozen peripheral blood mononuclear cells (PBMC) from lower-risk MDS patients (n=7) were treated with 1 \( \mu M \) lenalidomide for 24 h, before analysis of supernatant S100A9 concentration by ELISA. Results are expressed relative to untreated cells. (B) PBMC from lower-risk MDS patients (n=7) were treated with 1 \( \mu M \) lenalidomide 30 min prior to addition of lipopolysaccharide (LPS) (1 \( \mu g/mL \); 24 h after stimulation, TNF\( \alpha \) ELISA was performed on the supernatants. Results are expressed as a percentage relative to LPS treatment alone. *mean \( P<0.05 \).
0.164, \(P=0.006\), and between S100A9 and erythropoietin concentrations \((r=-0.148, P=0.01)\), whereas there was no discernible relationship between IL-1β and erythropoietin concentrations (Table 2A and Online Supplementary Figure S7). As expected, we also found significant positive correlations between the concentrations of the inflammatory protein S100A9 and TNFα \((r=0.294, P<0.001)\), S100A9 and IL-1β \((r=0.180, P=0.002)\), as well as IL-1β and TNFα \((r=0.262, P<0.001)\) (Table 2B). These findings support the notion that S100A9 and TNFα suppress renal erythropoietin elaboration and the endocrine response to anemia in non-del(5q) MDS.

**Relationships between inflammatory protein concentrations and response to treatment with erythropoietic agents**

Within the cohort of patients with non-del(5q) MDS, 159 were studied prior to ESA treatment and 159 prior to lenalidomide ± erythropoietin treatment (Table 1). ESA responders had significantly higher serum TNFα concentrations than non-responders \((8.57 \text{ pg/mL} \text{ versus} 8.79 \text{ pg/mL, respectively}; P=0.001)\) with a corresponding significantly lower erythropoietin concentration in ESA responders \((36 \text{ mU/mL} \text{ versus} 113 \text{ mU/mL, respectively}; P=0.0001)\). Erythroid response rate was 43% versus 55% in patients with low versus high TNFα concentration, respectively (Figure 4A). There was no significant relationship between S100A9 serum concentration and erythropoietin response (data not shown). Nevertheless, erythropoietin concentration was a better biomarker of response to ESA than was TNFα concentration even after adjustment for erythropoietin concentration. Finally, among patients treated with lenalidomide or lenalidomide ± erythropoietin, responding patients had significantly lower serum TNFα concentrations \((P=0.02)\) while there was no relationship with S100A9 concentration \((P=0.21)\). Considering responses to lenalidomide, we observed a significant difference in erythroid response rate \((62\% \text{ versus} 12\% \text{ for patients with low versus high S100A9 serum concentration, respectively}; P=0.03)\) (Figure 4B).

**Discussion**

Pro-inflammatory cytokines have long been implicated as key effectors of anemia in disorders of chronic inflamma-

**Table 2. (A) Correlations between concentrations of inflammatory proteins and erythropoietin (EPO) in patients’ serum. (B) Relationships between inflammatory proteins.**

<table>
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<th>Inflammatory parameters</th>
<th>EPO ((r; P\text{-value}))</th>
<th>S100A9 ((r; P\text{-value}))</th>
<th>TNFα ((r; P\text{-value}))</th>
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<td>S100A9</td>
<td>((-0.148; 0.01))</td>
<td>((-0.164; 0.006))</td>
<td>((-0.180; 0.002))</td>
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<tr>
<td>TNFα</td>
<td>((-0.148; 0.01))</td>
<td>((-0.148; 0.01))</td>
<td>((-0.180; 0.002))</td>
</tr>
<tr>
<td>IL-1β</td>
<td>((-0.164; 0.006))</td>
<td>((-0.180; 0.002))</td>
<td>((-0.180; 0.002))</td>
</tr>
</tbody>
</table>

Table 2. (A) Correlations between concentrations of inflammatory proteins and erythropoietin (EPO) in patients’ serum. (B) Relationships between inflammatory proteins.
higher serum TNFα concentrations. Together, these findings indicate that S100A9 and its transcriptional target, TNFα, directly suppress erythropoietin elaboration and endocrine response to anemia in MDS and may be useful biomarkers of response to treatment with lenalidomide or recombinant erythropoietin, meriting further investigation. More importantly, our findings suggest that therapeutic strategies that either neutralize or suppress S100A9 may improve erythropoiesis in lower-risk MDS by suppressing inflammatory cytokine generation and restoring endocrine erythropoietin response to anemia.

Acknowledgments
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References
20. Hori S, Nomura T, Sakaguchi S. Control of self-tolerance versus autoimmunity by the generation and restoring endocrine erythropoietin response to anemia in MDS. We found that the serum concentration of TNFα was significantly lower and serum concentration of S100A9 was higher in lenalidomide-responsive patients (P=0.03). Together, these findings indicate that S100A9 and its transcriptional target, TNFα, directly suppress erythropoietin elaboration and endocrine response to anemia in MDS and may be useful biomarkers of response to treatment with lenalidomide or recombinant erythropoietin, meriting further investigation. More importantly, our findings suggest that therapeutic strategies that either neutralize or suppress S100A9 may improve erythropoiesis in lower-risk MDS by suppressing inflammatory cytokine generation and restoring endocrine erythropoietin response to anemia.

Of NF-kB. Our findings support the notion that these inflammatory cytokines similarly suppress erythropoietin production in vivo in lower-risk MDS patients. Serum erythropoietin concentration was inversely related to S100A9 and TNFα concentrations. Furthermore, serum TNFα concentration was significantly higher in patients responding to treatment with recombinant erythropoietin than in non-responders (P=0.03). Previous investigations showed that higher serum TNFα concentration predicted resistance to ESA: our data do, therefore, need to be confirmed in a larger study. Of particular interest, lenalidomide suppressed nuclear translocation of NF-kB to mitigate the suppression of erythropoietin production in HepG2 cells by both S100A9 and TNFα. The ability of lenalidomide to modulate cytokine activation may not only reduce progenitor cell injury, but may also relieve repression of renal erythropoietin elaboration and, therefore, the endocrine response to anemia in MDS. We found that the serum concentration of TNFα was significantly lower and serum concentration of S100A9 was higher in lenalidomide-responsive patients (P=0.03). Together, these findings indicate that S100A9 and its transcriptional target, TNFα, directly suppress erythropoietin elaboration and endocrine response to anemia in MDS and may be useful biomarkers of response to treatment with lenalidomide or recombinant erythropoietin, meriting further investigation. More importantly, our findings suggest that therapeutic strategies that either neutralize or suppress S100A9 may improve erythropoiesis in lower-risk MDS by suppressing inflammatory cytokine generation and restoring endocrine erythropoietin response to anemia.