Increased Circulating CSF-1 (M-CSF) in Myeloproliferative Disease: Association With Myeloid Metaplasia and Peripheral Bone Marrow Extension

By Harriet S. Gilbert, Vincent Praloran, and E. Richard Stanley

Myeloproliferative disease (MPD) is heterogeneous in phenotypic expression and may display features consistent with expansion and activation of the monocyte/macrophage population during its course. The role of colony-stimulating factor-1 (CSF-1) in the pathophysiology of MPD was investigated by measuring circulating CSF-1 levels and examining their relationship to disease phenotype. Serum CSF-1 concentrations, measured by radioimmunoassay, were elevated in all MPD phenotypes. CSF-1 levels differed significantly between groups of patients with essential thrombocytemia, polycythemia vera, and postpolycythemic or agnogenic myeloid metaplasia (in ascending order). CSF-1 serum levels were positively correlated with spleen size and the degree of peripheral bone marrow extension, determined by scintigraphy using a macrophage-seeking isotope. There was no correlation between CSF-1 concentration and circulating levels of erythrocytes, neutrophils or platelets, or the presence of bone marrow fibrosis. Elevated serum CSF-1 levels appear to be associated with an expanded monocyte/macrophage population in MPD. In view of the known cooperativity between CSF-1 and other growth factors in regulating hematopoiesis, the finding of increased serum CSF-1 concentrations and its association with myeloid metaplasia and bone marrow extension may indicate a pathophysiologic role for CSF-1 in determining the phenotypic expression of MPD.

From the Departments of Medicine and Developmental Biology and Cancer, Albert Einstein College of Medicine, Bronx, NY.

Submitted November 28, 1988; accepted June 1, 1989.

Supported in part by Grants CA 31656 (to H.S.G.), CA 26504, CA 32551 (to E.R.S.) from the National Cancer Institute, and Grant RR-50 from the Division of Research Resources, US Public Health Service.

Address reprint requests to Harriet S. Gilbert, MD, Department of Medicine, Albert Einstein College of Medicine, 1300 Morris Park Ave, Bldg VE, Room 230, Bronx, NY 10461.

Dr Praloran's present address is Department of Hematology, Centre Hospitalier Regional de Nantes, Hotel Dieu BP 1005, 44035-Nantes Cedex, France.

The publication costs of this article were defrayed in part by page charge payment. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. section 1734 solely to indicate this fact.

© 1989 by Grune & Stratton, Inc.

0006-4971/89/7404-0036$3.00/0

Increased Circulating CSF-1 (M-CSF) in Myeloproliferative Disease: Association With Myeloid Metaplasia and Peripheral Bone Marrow Extension

By Harriet S. Gilbert, Vincent Praloran, and E. Richard Stanley

Myeloproliferative disease (MPD) is heterogeneous in phenotypic expression and may display features consistent with expansion and activation of the monocyte/macrophage population during its course. The role of colony-stimulating factor-1 (CSF-1) in the pathophysiology of MPD was investigated by measuring circulating CSF-1 levels and examining their relationship to disease phenotype. Serum CSF-1 concentrations, measured by radioimmunoassay, were elevated in all MPD phenotypes. CSF-1 levels differed significantly between groups of patients with essential thrombocytemia, polycythemia vera, and postpolycythemic or agnogenic myeloid metaplasia (in ascending order). CSF-1 serum levels were positively correlated with spleen size and the degree of peripheral bone marrow extension, determined by scintigraphy using a macrophage-seeking isotope. There was no correlation between CSF-1 concentration and circulating levels of erythrocytes, neutrophils or platelets, or the presence of bone marrow fibrosis. Elevated serum CSF-1 levels appear to be associated with an expanded monocyte/macrophage population in MPD. In view of the known cooperativity between CSF-1 and other growth factors in regulating hematopoiesis, the finding of increased serum CSF-1 concentrations and its association with myeloid metaplasia and bone marrow extension may indicate a pathophysiologic role for CSF-1 in determining the phenotypic expression of MPD.

From the Departments of Medicine and Developmental Biology and Cancer, Albert Einstein College of Medicine, Bronx, NY.

Submitted November 28, 1988; accepted June 1, 1989.

Supported in part by Grants CA 31656 (to H.S.G.), CA 26504, CA 32551 (to E.R.S.) from the National Cancer Institute, and Grant RR-50 from the Division of Research Resources, US Public Health Service.

Address reprint requests to Harriet S. Gilbert, MD, Department of Medicine, Albert Einstein College of Medicine, 1300 Morris Park Ave, Bldg VE, Room 230, Bronx, NY 10461.

Dr Praloran's present address is Department of Hematology, Centre Hospitalier Regional de Nantes, Hotel Dieu BP 1005, 44035-Nantes Cedex, France.

The publication costs of this article were defrayed in part by page charge payment. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. section 1734 solely to indicate this fact.

© 1989 by Grune & Stratton, Inc.

0006-4971/89/7404-0036$3.00/0

Increased Circulating CSF-1 (M-CSF) in Myeloproliferative Disease: Association With Myeloid Metaplasia and Peripheral Bone Marrow Extension

By Harriet S. Gilbert, Vincent Praloran, and E. Richard Stanley

Myeloproliferative disease (MPD) is heterogeneous in phenotypic expression and may display features consistent with expansion and activation of the monocyte/macrophage population during its course. The role of colony-stimulating factor-1 (CSF-1) in the pathophysiology of MPD was investigated by measuring circulating CSF-1 levels and examining their relationship to disease phenotype. Serum CSF-1 concentrations, measured by radioimmunoassay, were elevated in all MPD phenotypes. CSF-1 levels differed significantly between groups of patients with essential thrombocytemia, polycythemia vera, and postpolycythemic or agnogenic myeloid metaplasia (in ascending order). CSF-1 serum levels were positively correlated with spleen size and the degree of peripheral bone marrow extension, determined by scintigraphy using a macrophage-seeking isotope. There was no correlation between CSF-1 concentration and circulating levels of erythrocytes, neutrophils or platelets, or the presence of bone marrow fibrosis. Elevated serum CSF-1 levels appear to be associated with an expanded monocyte/macrophage population in MPD. In view of the known cooperativity between CSF-1 and other growth factors in regulating hematopoiesis, the finding of increased serum CSF-1 concentrations and its association with myeloid metaplasia and bone marrow extension may indicate a pathophysiologic role for CSF-1 in determining the phenotypic expression of MPD.

From the Departments of Medicine and Developmental Biology and Cancer, Albert Einstein College of Medicine, Bronx, NY.

Submitted November 28, 1988; accepted June 1, 1989.

Supported in part by Grants CA 31656 (to H.S.G.), CA 26504, CA 32551 (to E.R.S.) from the National Cancer Institute, and Grant RR-50 from the Division of Research Resources, US Public Health Service.

Address reprint requests to Harriet S. Gilbert, MD, Department of Medicine, Albert Einstein College of Medicine, 1300 Morris Park Ave, Bldg VE, Room 230, Bronx, NY 10461.

Dr Praloran's present address is Department of Hematology, Centre Hospitalier Regional de Nantes, Hotel Dieu BP 1005, 44035-Nantes Cedex, France.

The publication costs of this article were defrayed in part by page charge payment. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. section 1734 solely to indicate this fact.

© 1989 by Grune & Stratton, Inc.

0006-4971/89/7404-0036$3.00/0

Increased Circulating CSF-1 (M-CSF) in Myeloproliferative Disease: Association With Myeloid Metaplasia and Peripheral Bone Marrow Extension

By Harriet S. Gilbert, Vincent Praloran, and E. Richard Stanley

Myeloproliferative disease (MPD) is heterogeneous in phenotypic expression and may display features consistent with expansion and activation of the monocyte/macrophage population during its course. The role of colony-stimulating factor-1 (CSF-1) in the pathophysiology of MPD was investigated by measuring circulating CSF-1 levels and examining their relationship to disease phenotype. Serum CSF-1 concentrations, measured by radioimmunoassay, were elevated in all MPD phenotypes. CSF-1 levels differed significantly between groups of patients with essential thrombocytemia, polycythemia vera, and postpolycythemic or agnogenic myeloid metaplasia (in ascending order). CSF-1 serum levels were positively correlated with spleen size and the degree of peripheral bone marrow extension, determined by scintigraphy using a macrophage-seeking isotope. There was no correlation between CSF-1 concentration and circulating levels of erythrocytes, neutrophils or platelets, or the presence of bone marrow fibrosis. Elevated serum CSF-1 levels appear to be associated with an expanded monocyte/macrophage population in MPD. In view of the known cooperativity between CSF-1 and other growth factors in regulating hematopoiesis, the finding of increased serum CSF-1 concentrations and its association with myeloid metaplasia and bone marrow extension may indicate a pathophysiologic role for CSF-1 in determining the phenotypic expression of MPD.

From the Departments of Medicine and Developmental Biology and Cancer, Albert Einstein College of Medicine, Bronx, NY.

Submitted November 28, 1988; accepted June 1, 1989.

Supported in part by Grants CA 31656 (to H.S.G.), CA 26504, CA 32551 (to E.R.S.) from the National Cancer Institute, and Grant RR-50 from the Division of Research Resources, US Public Health Service.

Address reprint requests to Harriet S. Gilbert, MD, Department of Medicine, Albert Einstein College of Medicine, 1300 Morris Park Ave, Bldg VE, Room 230, Bronx, NY 10461.

Dr Praloran's present address is Department of Hematology, Centre Hospitalier Regional de Nantes, Hotel Dieu BP 1005, 44035-Nantes Cedex, France.

The publication costs of this article were defrayed in part by page charge payment. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. section 1734 solely to indicate this fact.

© 1989 by Grune & Stratton, Inc.

0006-4971/89/7404-0036$3.00/0
cells that express low levels of the CSF-1 receptor. An increase in circulating CSF-1 concentration might be expected to stimulate monocyte/macrophage differentiation, proliferation, and survival, thereby expanding and increasing the function of this population in MPD. Since CSF-1 is present in serum in concentrations that are detectable by radioimmunoassay, this hypothesis was tested by measuring circulating CSF-1 levels in MPD and determining their relation to disease phenotype.

MATERIALS AND METHODS

Patient material. Serum was obtained from blood samples drawn with informed consent from age-matched, hematologically normal subjects and patients with MPD who met the diagnostic criteria of the Polycythemia Vera Study Group. The protocol was approved by the local Committee for Clinical Investigation. Disease activity was evaluated by a comprehensive medical history, physical examination, complete blood count, serum chemistry determinations, bone marrow biopsy, serum unsaturated vitamin B12 binding capacity, and liver, spleen, and bone marrow imaging with 99mTc-SC. Patients were categorized by MPD phenotype as essential thrombocythemia (ET), polycythemia vera (PV), postpolycythemic myeloid metaplasia (PPMyM), and agnogenic myeloid metaplasia (AMyM). Subjects were free from clinically evident infection and were not receiving myelosuppressive treatment at the time of study, except for two patients with PV and one with AMyM, who were being treated with hydroxyurea. Bone marrow biopsies were examined for the presence of collagen fibrosis according to published criteria.17

CSF-1 assay. Serum CSF-1 was measured using a modification of a previously published radioimmunoassay (RIA). Modifications improved the sensitivity of the assay and included slight changes in the pH and protein concentration of the buffers and a change in the technique used for precipitation of the CSF-1 antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immunization with purified human urinary CSF-1. Normal rabbit serum (NRS) diluted 1/10 in RIA buffer, pH 7.5, was used for the preparation of the antigen-antibody complexes. Purified human CSF-1 from medium conditioned by the M1A PaCa cell line or recombinant CSF-1 was obtained from Cetus Corp (Emeryville, CA) and iodinated as described. Anti-human CSF-1 antiserum was raised in rabbits by immu

Fig 1. Serum CSF-1 concentrations in hematologically normal subjects and patients with myeloproliferative disease, grouped by syndrome. Dots represent actual values. Boxes characterize the population. The center line of the box is the median that splits the population in half. The edges of the box are the hinges that split the halves in half again and demarcate the 25th to 75th percentile of the population. Outliers with values falling beyond 1.5 time the range between the hinges are indicated by an asterisk. The vertical lines extend from the hinges to the largest or smallest nonoutlier value.27 Intergroup comparisons by Wilcoxon signed ranks testing showed significantly higher CSF-1 values in the MPD syndromes (P values shown in Table 1). Significant intergroup differences were observed in serum CSF concentrations amongst the ET, PV, and the combined PPMym and AMym groups, ranked in ascending order.
INCREASED CSF-1 IN MYELOPROLIFERATIVE DISEASE

Table 1. Serum CSF-1 Concentrations (ng/mL) in Hematologically Normal Subjects and Patients With Myeloproliferative Syndromes

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>N</th>
<th>Mean ± SD</th>
<th>Median</th>
<th>Range</th>
<th>P value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>34</td>
<td>4.9 ± 1.1</td>
<td>4.8</td>
<td>1.7-7.1</td>
<td>.04</td>
</tr>
<tr>
<td>ET</td>
<td>9</td>
<td>5.9 ± 1.3</td>
<td>6.0</td>
<td>3.4-8.1</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>PV</td>
<td>23</td>
<td>7.4 ± 1.8</td>
<td>7.4</td>
<td>3.6-10.7</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>PPMym</td>
<td>7</td>
<td>12.4 ± 3.3</td>
<td>10.8</td>
<td>9.0-18.2</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>AMYM</td>
<td>12</td>
<td>10.0 ± 7.2</td>
<td>10.2</td>
<td>6.5-31.6</td>
<td>&lt;.005</td>
</tr>
</tbody>
</table>

*Compared with Normal group.
†Population normally distributed.
‡Population not normally distributed.

lower extremities as grade 1 (normal; no scintigraphic activity below the upper third of the femur); grade 2 (expansion to femur and knee; grade 3 (expansion to femur and tibia; scintigraphic activity extending to the lower femur and tibia); grade 4 (expansion involving the entire lower extremity). MPD patients with bone marrow expansion involving the entire lower extremity had significantly higher levels of CSF-1 than those without peripheral marrow expansion and those with marrow expansion limited to the femur (Fig 2). Since the upper third of the femur normally contains some hematopoietic tissue, when considered in terms of bone marrow mass, extension of bone marrow to the femur and tibia would represent a significantly greater increment in hematopoietic tissue than would expansion only to the middle and lower third of the femur. If bone marrow expansion and CSF-1 concentrations are related, the observed significant increase in CSF-1 levels in patients with expansion involving the entire lower extremities would be expected.

Since fibroblasts are a source of CSF-1 and reactive fibroblast proliferation is a complication of MPD often associated with myeloid metaplasia, the presence of collagen fibrosis was determined from bone marrow biopsies of the posterior iliac crest and the relation of serum CSF-1 concentrations to the presence of fibrosis in the 17 patients with myeloid metaplasia was examined. There was no significant difference in CSF-1 concentrations between patients in the myeloid metaplasia group with (n = 9) and without (n = 8) bone marrow fibrosis (mean ± SD, 11.4 ± 4.3 v 12.9 ± 7 ng/mL, respectively; P > .05).

The lack of correlation between serum CSF-1 concentrations and leukocyte counts in the MPD population is notable and is in agreement with our findings in four patients with leukemia (two acute myeloblastic, one acute lymphoblastic, one chronic myelocytic) before and during intensive chemotherapy. All had normal CSF-1 levels before therapy and showed no significant change during treatment (Praloran V, Gilbert HS, Stanley ER, unpublished observations). Hanamura et al reported elevated serum M-CSF levels in one half of a group of patients with hematologic malignancy after anticancer chemotherapy, and postulated that this was a response to neutropenia. The persistence of normal M-CSF levels in the other half was attributed to significant damage to CSF-producing cells by intensive chemotherapy. Serum M-CSF was also elevated in patients with infection and varying degrees of neutrophilia, as well as in pregnant women with normal neutrophil counts.

Elevated serum CSF-1 could result from increased production and/or decreased catabolism. The effects and turnover of circulating CSF-1 are mediated by the CSF-1 receptor present on the target populations. This receptor is encoded by the c-fms protooncogene and expressed selectively in normal mononuclear phagocytes. The cellular source of circulating CSF-1 in humans is poorly understood. While monocytes express low levels of CSF-1 transcripts, monocyte activation by phorbol ester induces CSF-1 expression. This suggests that under certain conditions monocytes may have the capacity to autostimulate certain functions through the production of CSF-1. Treatment of resting human monocytes with GM-CSF similarly induces CSF-1 expression. Induction of CSF-1 transcripts in monocytes is associated with synthesis of the CSF-1 gene product. This induction occurred during the continued expression of c-fms transcripts, although there was a partial down-regulation of c-fms transcription. Other cytokines, including T-cell-derived γ-interferon and interleukin-3, appear capable of inducing CSF-1 in monocytes. Based on the behavior of normal human monocytes, a hypothesis to explain increased circulating concentrations of CSF-1 in MPD would be the presence of an expanded and activated monocyte/macrophage population in which CSF-1 production is increased, either by an autocrine mechanism or by induction by other HGFs, and CSF-1 receptor (c-fms) expression is down-regulated, leading to decreased removal and catabolism of circulating CSF-1. The alterations in the metabolism of CSF-1 and other hematopoietic growth factors and cytokines...
that might alter CSF-1 metabolism in MPD remain to be investigated.

The elevated serum CSF-1 in MPD could result from either increased release into or decreased clearance from the circulation, or both. The cellular and tissue sources of circulating CSF-1 are not known and may vary in different imposed situations. However, in mice, clearance of circulating CSF-1 occurs by CSF-1 receptor-mediated endocytosis and intracellular destruction by macrophages of the liver and spleen. Elevated plasma CSF-1 concentrations observed in MPD could, therefore, result from a decrease in the number and/or function of these macrophages. Whatever the cause of increased CSF-1 levels in MPD, their association with myeloid metaplasia and bone marrow expansion suggests that CSF-1 may affect the amount and distribution of hematopoietically active tissue in MPD by direct stimulation of the macrophage population and/or by stimulation of the PHPC and its other progeny through cooperation with other HGFs. The availability of sufficient quantities of recombinant human M-CSF for human administration will facilitate studies of the biology of CSF-1 in MPD to test these hypotheses and obtain insights into potential clinical applications of CSF-1 for manipulation of the hematopoietic organ.

ACKNOWLEDGMENT

We thank Claudia Saez and Anne Palestrone for excellent technical assistance.

REFERENCES

Increased circulating CSF-1 (M-CSF) in myeloproliferative disease: association with myeloid metaplasia and peripheral bone marrow extension

HS Gilbert, V Praloran and ER Stanley