Effect of Hydroxyurea on Mortality and Morbidity in Adult Sickle Cell Anemia
Risks and Benefits Up to 9 Years of Treatment

Martin H. Steinberg, MD
Franca Barton, MS
Oswaldo Castro, MD
Charles H. Pegelow, MD†
Samir K. Ballas, MD
Abdullah Kutlar, MD
Eugene Orringer, MD
Rita Bellevue, MD
Nancy Olivieri, MD
James Eckman, MD
Mala Varma, MD
Gloria Ramirez, MD
Brian Adler, MD
Wally Smith, MD
Timothy Carlos, MD
Kenneth Ataga, MD
Laura DeCastro, MD
Carolyn Bigelow, MD
Yogen Saunthararajah, MD
Margaret Telfer, MD
Elliot Vichinsky, MD
Susan Claster, MD
Myron Waclawiw, PhD
Duane Bonds, MD
Michael Terrin, MD, MPH

Context  Hydroxyurea increases levels of fetal hemoglobin (HbF) and decreases morbidity from vaso-occlusive complications in patients with sickle cell anemia (SCA). High HbF levels reduce morbidity and mortality.

Objective  To determine whether hydroxyurea attenuates mortality in patients with SCA.

Design  Long-term observational follow-up study of mortality in patients with SCA who originally participated in the randomized, double-blind, placebo-controlled Multicenter Study of Hydroxyurea in Sickle Cell Anemia (MSH), conducted in 1992-1995, to determine if hydroxyurea reduces vaso-occlusive events. In the MSH Patients’ Follow-up, conducted in 1996-2001, patients could continue, stop, or start hydroxyurea. Data were collected during the trial and in the follow-up period.

Setting  Inpatients and outpatients in 21 sickle cell referral centers in the United States and Canada.

Patients  Two-hundred ninety-nine adult patients with frequent painful episodes enrolled in the follow-up. Follow-up data through May 2001 were complete for 233 patients.

Intervention  In the MSH, patients were randomly assigned to receive hydroxyurea (n = 152) or placebo (n = 147).

Main Outcome Measures  Mortality, HbF levels, painful episodes, acute chest syndrome, and blood cell counts. The randomized trial was not designed to detect specified differences in mortality.

Results  Seventy-five of the original 299 patients died, 28% from pulmonary disease. Patients with reticulocyte counts less than 250000/mm³ and hemoglobin levels lower than 9 g/dL had increased mortality (P = .002). Cumulative mortality at 9 years was 28% when HbF levels were lower than 0.5 g/dL after the trial was completed compared with 15% when HbF levels were 0.5 g/dL or higher (P = .03). Individuals who had acute chest syndrome during the trial had 32% mortality compared with 18% of individuals without acute chest syndrome (P = .02). Patients with 3 or more painful episodes per year during the trial had 27% mortality compared with 17% of patients with less frequent episodes (P = .06). Taking hydroxyurea was associated with a 40% reduction in mortality (P = .04) in this observational follow-up with self-selected treatment. There were 3 cases of cancer, 1 fatal.

Conclusions  Adult patients taking hydroxyurea for frequent painful sickle cell episodes appear to have reduced mortality after 9 years follow-up. Survival was related to HbF levels and frequency of vaso-occlusive events. Whether indications for hydroxyurea treatment should be expanded is unknown.

JAMA. 2003;289:1645-1651

©2003 American Medical Association. All rights reserved.
Multicenter Study of Hydroxyurea in Sickle Cell Anemia (MSH), showed that, over 2.5 years, hydroxyurea diminished the morbidity of SCA in adults with frequent painful episodes by reducing the incidence of painful episodes and acute chest syndrome by nearly half.2,3 Hydroxyurea increased fetal hemoglobin (HbF) concentration while hemolysis was diminished and neutrophil counts fell.1,3 In SCA, HbF levels are inversely related to mortality.1 Decreased morbidity due to hydroxyurea could be associated with reduced mortality. Following the completion of the MSH, patients enrolled in an observational follow-up study, selecting with their physicians whether to continue, start, or stop treatment with hydroxyurea. Follow-up focused on detecting potential complications of treatment such as neoplasia and cerebrovascular disease, and on ascertaining the causes of death. Cumulative mortality was analyzed according to exposure to hydroxyurea and data from the MSH on acute chest syndrome, acute painful episodes, HbF levels, reticulocyte counts, and neutrophil counts, assessed both before randomization and at the conclusion of the trial.

METHODS

The MSH Patients’ Follow-up is an observational study of inpatients and outpatients in 21 sickle cell referral centers in the United States and Canada who participated from 1992 to 1995 in the MSH to test whether hydroxyurea could reduce the number of vaso-occlusive events in adults with moderate to severe SCA.5 Requirements for MSH entry included a history of more than 3 painful episodes in the 12 months prior to enrollment and pretreatment hemoglobin (HbF) levels, reticulocyte counts, and neutrophil counts, assessed both before randomization and at the conclusion of the trial. (most patients were seen more often for medical care) when data on hydroxyurea use and reportable events including stroke, renal failure, hepatic failure, cancer, and sepsis were collected. Fatal events and serious illnesses were documented from medical records. Central review of medical records, and autopsy results when these were available, were completed to establish causes of death. Patients provided internal review board–approved written informed consent for enrollment in the follow-up study.

Laboratory Methods

Methods for blood cell counts and measurement of HbF levels have been previously reported.2,3,5,6 A single pretreatment value for each laboratory determination was computed by averaging all pretreatment values of that measurement for each patient. A single 2-year value was computed by averaging available data during the final 18 to 24 months of study treatment.2 Levels of HbF were expressed as absolute levels of HbF in g/dL. HbF (g/dL] = HbF [%] × hemoglobin concentration [g/dL]) because no biological mechanism of HbF production exists that would regulate HbF as a percentage of total hemoglobin.

Statistical Methods

Patients were grouped for comparisons in a variety of ways, including the treatment group (hydroxyurea or placebo) to which they were originally assigned, and according to characteristics measured later during the MSH, such as HbF level and painful crisis rate at 2 years. For discrete variables, χ² tests were used to compare the frequency of patient characteristics.7 Continuous measurements are presented as mean (SD) and compared using analysis of variance.8

Mortality in 3-month intervals was assessed using general estimating equation logistic models based on hydroxyurea usage (yes or no) during each interval.9 Analyses were performed using SAS v7.6 (SAS Institute Inc, Cary, NC). Mortality is also reported as cumulative 9-year event rate from Kaplan-Meier survival curves.10 Differences in mortality between patients randomly assigned to receive hydroxyurea and those assigned to receive placebo (ie, intention-to-treat analyses) and between post hoc groups (eg, HbF levels at 2 years) were tested with the log-rank statistic.10 Patients were censored at the time they were last documented to be alive by clinical center staff. Hydroxyurea usage was measured as taken, not taken, or unknown for each month during the trial and follow-up period until the patient died or was censored, and was summarized as total months of taking hydroxyurea (never, <1 year, or ≥1 year).

The MSH was not designed to detect specified differences in mortality. Because of multiple testing that has occurred in secondary analyses of data from these patients, the strength of statistical evidence for differences between the original treatment groups was predetermined at a more stringent level than the primary end point trial design of α = .05.3 Observed differences carrying a nominal P value of less than .05 should not be taken as conclusive evidence of a difference. Because inference is not formal with the comparisons presented now—most of which are observational and not randomized—the nominal P values should be regarded as indicators of association, not tests of a priori hypotheses, and should be interpreted cautiously.

RESULTS

Follow-up data for up to 9 years through May 2001 were complete for 233 (77.9%) of the 299 patients enrolled in the long-term follow-up. Eight patients died during the randomized trial (2 receiving hydroxyurea, 6 receiving placebo). Hydroxyurea use in follow-up is unknown for 13 patients (4.3%) originally randomly assigned to receive hydroxyurea and for 15 patients (5.0%) assigned to receive placebo. Vital status was not entered into our database for 27 patients (9.0%).

Many patients were treated with hydroxyurea for at least 1 year since the start of the trial (TABLE 1). Ninety-six (32%) patients never received hydroxy-
urea; 48 (16%) received hydroxyurea for less than 1 year and 156 (52%) received hydroxyurea for 1 or more years. Twenty-five percent of patients (n=75) who volunteered for the MSH died during the trial or follow-up, reflecting the severity of disease, which was a criterion for enrollment, and the high death rate in adult SCA.1 Twenty-eight percent of these deaths were due to pulmonary complications (TABLE 2).

When analyzed according to the original assignment (regardless of the patient’s choice of treatment after the randomized phase of the study was completed), mortality was similar in the original 2 treatment groups (P = .35; FIGURE, A). Cumulative mortality analyzed according to clinical events and laboratory measurements at the conclusion of randomized treatment is shown in the Figure (panels B-F). Of 276 patients who had HbF levels measured approximately 2 years into the MSH, patients with HbF levels lower than 0.5 g/dL had a 28% cumulative mortality through 9 years compared with 15% mortality in patients whose HbF levels were 0.5 g/dL or higher (P = .03 by log-rank test; Figure, B). The effect on mortality was similar when HbF levels lower than 0.75 g/dL and those 0.75 g/dL or higher were analyzed. Cumulative mortality through 9 years in the 199 patients with pretreatment HbF levels lower than 0.5 g/dL was 32% compared with 15% mortality in patients whose pretreatment HbF levels were 0.5 g/dL or higher (P = .01). Before treatment, 68% of patients randomly assigned to receive hydroxyurea and 65% of those assigned to receive placebo had HbF levels of lower than 0.5 g/dL. When the trial was completed, 38% of the patients assigned to receive hydroxyurea with initial HbF levels lower than 0.5 g/dL had HbF levels 0.5 g/dL or higher. In contrast, only 8% of patients receiving placebo with baseline HbF levels lower than 0.5 g/dL had final HbF levels 0.5 g/dL or higher.

Neutrophil counts before random assignment and treatment were not associated with mortality. Mortality increased with the number of transfusions in the year before enrollment (P = .001). The proportion of patients with a peak hemoglobin F level of less than 1% was higher among patients who died during follow-up than among those who were alive (22% vs 11%; P = .05). Overall, more than half (54%) of the patients had a peak HbF level of less than 1%. No deaths were thought to be treatment related.

Table 1. Cumulative Event Rates During MSH Follow-up (9 Years)

<table>
<thead>
<tr>
<th>Treatment Assigned in MSH</th>
<th>Time Receiving Hydroxyurea in MSH and Follow-up</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Total Hydroxyurea Placebo Never /H11021 /H11350</td>
</tr>
<tr>
<td>Total, No.</td>
<td>299 152 147 47 33 219</td>
</tr>
<tr>
<td>Person-years†</td>
<td>2264 1174 1091 314 219 1731</td>
</tr>
<tr>
<td>Follow-up, mean (SD)</td>
<td>7.6 (2.0) 7.7 (1.7) 7.4 (2.1) 6.7 (2.9) 6.6 (2.7) 7.8 (1.4)</td>
</tr>
<tr>
<td>Events</td>
<td></td>
</tr>
<tr>
<td>Death, No. (%)</td>
<td>75 (25.1) 36 (23.7) 39 (26.5) 14 (29.8) 15 (45.5) 46 (21.0)</td>
</tr>
<tr>
<td>No. per 100 person-years</td>
<td>3.3 3.1 3.6 4.5 6.8 2.7</td>
</tr>
<tr>
<td>Stroke, No. (%)</td>
<td>14 (4.7) 8 (5.3) 6 (4.1) 1 (2.1) 2 (6.1) 11 (5.0)</td>
</tr>
<tr>
<td>No. per 100 person-years</td>
<td>0.6 0.7 0.5 0.3 0.9 0.6</td>
</tr>
<tr>
<td>Renal failure, No. (%)</td>
<td>28 (9.4) 14 (9.2) 14 (9.5) 9 (19.1) 3 (9.1) 16 (7.3)</td>
</tr>
<tr>
<td>No. per 100 person-years</td>
<td>1.2 1.2 1.3 2.9 1.4 0.9</td>
</tr>
<tr>
<td>Hepatic failure, No. (%)</td>
<td>13 (4.3) 3 (2.0) 10 (6.8) 5 (10.6) 2 (6.1) 6 (2.7)</td>
</tr>
<tr>
<td>No. per 100 person-years</td>
<td>0.6 0.3 0.9 1.6 0.9 0.3</td>
</tr>
<tr>
<td>Malignancy, No. (%)‡</td>
<td>2 (0.7) 1 (0.7) 1 (0.7) 0 0 2 (0.9)</td>
</tr>
<tr>
<td>No. per 100 person-years</td>
<td>0.1 0.1 0.1 0 0 0.1</td>
</tr>
<tr>
<td>Sepsis/infection, No. (%)</td>
<td>38 (12.7) 18 (11.8) 20 (13.6) 3 (6.4) 6 (18.2) 29 (13.2)</td>
</tr>
<tr>
<td>No. per 100 person-years</td>
<td>1.7 1.5 1.8 1.0 2.7 1.7</td>
</tr>
</tbody>
</table>

Abbreviation: MSH, Multicenter Study of Hydroxyurea in Sickle Cell Anemia.
*Includes 6 patients who took hydroxyurea for less than 3 months, counted as 0 months receiving hydroxyurea in Table 4.
†From start of the MSH to patient death or data cutoff.
‡A third malignancy in the MSH Patients’ Follow-up occurred after data files were closed for this analysis and is not included in this table.

Table 2. Cause of Death According to Original Treatment Assignment in Patients in the MSH

<table>
<thead>
<tr>
<th>Causes of Death*</th>
<th>Original Treatment Assignment</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Hydroxyurea, No. (%) (n = 152)</td>
</tr>
<tr>
<td>Total deaths</td>
<td>36 (23.7)</td>
</tr>
<tr>
<td>Pulmonary disease</td>
<td>7 (4.6)</td>
</tr>
<tr>
<td>Death during crisis</td>
<td>4 (2.6)</td>
</tr>
<tr>
<td>Sepsis/infection</td>
<td>2 (1.3)</td>
</tr>
<tr>
<td>Cardiovascular disease</td>
<td>1 (0.7)</td>
</tr>
<tr>
<td>Cerebrovascular</td>
<td>6 (3.9)</td>
</tr>
<tr>
<td>Hepatic disease</td>
<td>2 (1.3)</td>
</tr>
<tr>
<td>Gastrointestinal disease</td>
<td>2 (1.3)</td>
</tr>
<tr>
<td>Renal disease</td>
<td>1 (0.7)</td>
</tr>
<tr>
<td>Unintentional injury/homicide</td>
<td>3 (2.0)</td>
</tr>
<tr>
<td>Undetermined</td>
<td>4 (2.6)</td>
</tr>
<tr>
<td>Not yet classified</td>
<td>4 (2.6)</td>
</tr>
</tbody>
</table>

Abbreviation: MSH, Multicenter Study of Hydroxyurea in Sickle Cell Anemia.
*The single death from malignancy in the MSH Patients’ Follow-up occurred after data files were closed for this analysis and is not included in this table.
associated with differences in mortality. Neutrophil counts above or below 5000/mm³ at the end of the clinical trial also did not predict mortality (P = .70 by log-rank test; Figure, C), although aggressive titration with hydroxyurea affects neutrophil counts. Similarly, neutrophil counts above or below 7000/mm³ did not affect mortality. Reticulocyte counts before randomization did not predict mortality. Patients with absolute reticulocyte counts less than 250000/mm³ after 2 years of treatment had a cumulative mortality of 37% compared with 18% for individuals with 250000/mm³ or more (P = .001). A subgroup of 63 patients with reticulocyte counts less than 250000/mm³ and hemoglobin concentrations lower than 9 g/dL had increased cumulative mortality up to 9 years of observation (38% dead at 9 years, P = .002 by log-rank test; Figure, D) They also had lower HbF levels (0.42 g/dL vs 0.62 g/dL; P = .005), higher mean (SD) serum creatinine levels (1.49 [1.77] vs 0.96 [0.23] mg/dL [131.7 [156.5] vs 84.9 [20.3] µmol/L]; P = .02) and received lower doses of hydroxyurea compared with other patient groups (Table 3).

Patients with no episodes of acute chest syndrome during the trial had a mortality of 18% compared with 32% in patients who had 1 or more episodes (P = .02; Figure, E). Individuals with fewer than 3 annual painful episodes during the clinical trial had a mortality of 17% compared with 27% in patients with 3 or more episodes annually (P = .06; Figure 1F).

As shown in Table 4, total months of exposure to hydroxyurea were related to original treatment assignment, duration of survival in follow-up, and choice of treatment in follow-up. By analyzing mortality in 3-month intervals according to hydroxyurea usage in the interval, death rates were reduced 40%
Two patients received placebo and then switched to hydroxyurea during 3-month intervals when patients were taking hydroxyurea [2.6 (5.8) deaths per 3-month period vs 1.5 (7.9) deaths per 3 months; \( P = .04 \)].

Three patients developed cancer. One individual who was randomly assigned to receive placebo, but who subsequently received hydroxyurea for 63 months, had carcinoma in situ of the uterine cervix at 7 years after randomization. A patient randomly assigned to hydroxyurea had 47 months of exposure to hydroxyurea in 3 periods: for less than 1 year during the MSH, after the clinical trial, and after mastectomies 3 years later. She had a history of fibrocystic disease of the breast and had had prophylactic bilateral mastectomies. Histological examination showed multifocal carcinoma in situ in the left breast. Both patients died of causes unrelated to carcinoma. A third patient randomly assigned to hydroxyurea who received long-term treatment with hydroxyurea [2.6 (5.8) deaths per 3-month period vs 1.5 (7.9) deaths per 3 months; \( P = .04 \)].

The MSH was designed to test whether hydroxyurea reduced the incidence of vaso-occlusive episodes in SCA and not to detect differences in mortality between treatment groups. However, our follow-up studies suggest that adults with moderate to severe SCA who take hydroxyurea have reduced mortality compared with patients not taking this drug. Our estimated overall reduction in mortality up to 9 years of observation is 40%. These results must be interpreted cautiously. Comparisons of patients receiving and not receiving hydroxyurea were no longer randomized after the initial treatment period of the MSH, so patients who might receive more medical care and better follow-up while taking hydroxyurea might live longer for reasons other than treatment.

### Table 4. Total Months That All Patients Randomly Assigned in the MSH Actually Received Hydroxyurea

<table>
<thead>
<tr>
<th>Receiving Hydroxyurea, mo</th>
<th>Total</th>
<th>Randomized to Receive Placebo, No./Deaths</th>
<th>Randomized to Receive Hydroxyurea</th>
<th>Hydroxyurea Use in Follow-up Period Unknown or Incompletely Documented</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Never Received Hydroxyurea</td>
<td>Subsequently Received Hydroxyurea</td>
<td>Stopped Taking Hydroxyurea</td>
</tr>
<tr>
<td>&lt;3†</td>
<td>53/15</td>
<td>36/13</td>
<td>...</td>
<td>1/0</td>
</tr>
<tr>
<td>3-11</td>
<td>27/14</td>
<td>...</td>
<td>19/9</td>
<td>6/5</td>
</tr>
<tr>
<td>12-17</td>
<td>13/5</td>
<td>...</td>
<td>9/4</td>
<td>3/0</td>
</tr>
<tr>
<td>18-23</td>
<td>10/4</td>
<td>...</td>
<td>4/1</td>
<td>5/3</td>
</tr>
<tr>
<td>24-36</td>
<td>48/14</td>
<td>...</td>
<td>15/4</td>
<td>17/4</td>
</tr>
<tr>
<td>36-45</td>
<td>31/8</td>
<td>...</td>
<td>13/3</td>
<td>1/0</td>
</tr>
<tr>
<td>48-59</td>
<td>23/6</td>
<td>...</td>
<td>12/2</td>
<td>...</td>
</tr>
<tr>
<td>60-71</td>
<td>32/3</td>
<td>...</td>
<td>22/1</td>
<td>...</td>
</tr>
<tr>
<td>72-83</td>
<td>22/6</td>
<td>...</td>
<td>2/0</td>
<td>...</td>
</tr>
<tr>
<td>84-95</td>
<td>17/0</td>
<td>...</td>
<td>...</td>
<td>...</td>
</tr>
<tr>
<td>96-107</td>
<td>23/1</td>
<td>...</td>
<td>...</td>
<td>...</td>
</tr>
<tr>
<td>Total</td>
<td>299/75</td>
<td>36/13</td>
<td>96/24</td>
<td>33/12</td>
</tr>
</tbody>
</table>

*Ellipses indicate not applicable.
†Counted as 0.

©2003 American Medical Association. All rights reserved.

(Reprinted) JAMA, April 2, 2003—Vol 289, No. 13 1649
HYDROXYUREA AND ADULT SICKLE CELL ANEMIA

April 2, 2003—Vol 289, No. 13 (Reprinted)

©2003 American Medical Association. All rights reserved.

ment. Moreover, assessment of the effect of hydroxyurea on mortality in an observational study is complex. The ability to take hydroxyurea (a predictor) is dependent on being alive (an outcome). Also, patients in the MSH were not typical of all adults with SCA; their average age was 32 years and they were selected for the trial because of frequent painful episodes, a predictor of mortality. However, it is compelling that patients who had marked clinical benefit, reduced crises, reduced chest syndrome, and a good bone marrow reserve after 2 years of receiving hydroxyurea as randomized were the ones with reduced mortality in follow-up (Figure). This, coupled with the more direct, albeit observational, comparison of 3 months receiving hydroxyurea vs 3 months not receiving the drug is consistent with a long-term salutary effect of hydroxyurea on mortality.

How hydroxyurea affects morbidity and mortality in young children or individuals with less severe disease is unknown. Clinical trials of hydroxyurea have not been conducted in patients with HbSC disease, another clinically important sickle hemoglobinopathy. There were no appreciable differences for mortality or the clinical events reported here in follow-up between the 2 initial treatment groups. The closest test of whether hydroxyurea has an impact on mortality in SCA would be a controlled study in children that could provide definitive data on long-term mortality effects. However, the death rate in children is lower than in older adults. Since hydroxyurea is likely to have similar short-term beneficial effects in children and adults, prolonged follow-up of a group of children receiving placebo may not be acceptable to families. In 93 patients with SCA, 87 of whom were younger than 20 years at the start of treatment, no severe adverse effects of treatment were found after a median of 3.5 years of observation.

Hydroxyurea reduces some vasoocclusive complications of SCA. While its mechanism of action is incompletely understood, hydroxyurea is associated with increased levels of HbF in SCA; increased levels of HbF are known to improve survival. In the MSH and other studies, hydroxyurea reduced the incidence of acute chest syndrome and acute painful episodes, events associated with increased mortality and influenced by HbF concentrations. Levels of HbF 0.5 g/dL or higher, absence of acute chest syndrome, and fewer than 3 painful episodes annually during the trial were each associated with nearly a 50% reduction in mortality. Almost half of the classified deaths in our study were due to pulmonary disease or sudden death during a painful episode. Our observation of reduced mortality associated with hydroxyurea is consistent with hydroxyurea reducing the incidence of these morbid events by nearly half in the MSH. These observations also confirm the link of HbF to mortality in sickle cell disease and suggest that the ability of hydroxyurea to increase levels of HbF may be associated with reduced mortality.

Besides HbF, proposed mediators of the clinical effectiveness of hydroxyurea include reductions in adherent reticulocytes that may help initiate vasoocclusion and in leukocytes that might release proinflammatory cytokines. Neutrophil counts have been associated with mortality in other diseases besides sickle cell disease. Leukocytosis is a predictor of the extent of and sudden death from coronary artery disease. Some beneficial effects of hydroxyurea in SCA have been associated with reduction in neutrophil counts during treatment. We found no relationship between decrements in neutrophil counts—also a predictor of HbF concentration in patients treated with hydroxyurea—and mortality, but the Cooperative Study of Sickle Cell Disease (CSSCD) found that lower leukocyte counts were associated with longevity. In the MSH, hydroxyurea was titrated to maximum tolerated doses that most often were the largest doses that could be taken without neutropenia. Consequently, most patients taking hydroxyurea developed some level of potentially dangerous neutropenia. Since hydroxyurea was associated with a strong beneficial effect on painful crisis frequency in the MSH, this design forced an association between neutropenia and painful crisis frequency—low white blood cell counts being associated with fewer crises. Since neutrophils are acute-phase reactants, the association in the MSH is in the same direction as the association of neutrophil counts with mortality in the CSSCD, but for a different reason. In the follow-up study, doses of hydroxyurea used by the patients’ physicians were less than the maximum tolerated doses in the MSH. During the follow-up, hydroxyurea effects, like increases in concentrations of HbF and total hemoglobin and decreases in bilirubin levels, were observed without notable effects on leukocyte counts (F.B., unpublished data). Although induction of neutropenia may play some role in the beneficial effects of hydroxyurea for SCA, the follow-up study data suggest that neutropenia is not a main mechanism of action.

Patients with reticulocyte counts less than 250 000/mm³ and hemoglobin concentrations lower than 9 g/dL had increased mortality. These individuals also had lower HbF levels, higher serum creatinine levels, and received lower doses of hydroxyurea. This combination of findings suggests that these patients had more severe disease and perhaps reduced marrow reserve and that, as a result, they were unable to tolerate sufficient hydroxyurea treatment to increase HbF levels. Anemia and reticulocytopenia may also be indications of early renal failure that is associated with a poor prognosis.

During the 9-year observation period we found little risk associated with the careful use of hydroxyurea in SCA. Yet, hydroxyurea must be taken indefinitely to be effective and is potentially mutagenic and carcinogenic. Other investigators have reported that 5% to 10% of patients with polycythemia vera and essential thrombocythemia, both preneoplastic myeloproliferative diseases, who received hydroxyurea developed acute leukemia. Whether or not this risk can be extrapolated to patients with SCA is uncertain. Three reports of patients with SCA treated with hydroxy-
urea who developed acute leukemia have been published, but the total number of patients treated is unknown.20-31 Our data suggest that the risk of leukemia in patients with SCA treated with hydroxyurea is much less than that observed in myeloproliferative disorders and that the risk of death from the complications of sickle cell disease is at least 10 times greater than the incidence of leukemia in these patients.

Intracerebral hemorrhage, the main cause of stroke in adult SCA, occurs at a rate of about 1 per 400 patient-years.32,33 Given our number of patients under observation, we should have been able to detect a doubling in the incidence of intracerebral hemorrhage, but no association of stroke with hydroxyurea was found. Cerebrovascular mortality occurred more frequently among patients originally assigned to hydroxyurea than placebo, but was not related to use of hydroxyurea in follow-up.

Hydroxyurea increased HbF concentrations and diminished vaso-occlusive complications in adults with moderate to severe SCA and these effects were associated with decreased mortality. After up to 9 years of follow-up, unexpected severe adverse effects of this treatment were not observed. Whether hydroxyurea should be given to patients with SCA and fewer vaso-occlusive events or to patients with HbSC disease remains to be determined. We conclude that underlying disease severity remains critical to determining the prognosis of adult SCA, but hydroxyurea may mitigate disease severity.

Author Affiliations: Boston University School of Medicine, Center of Excellence in Sickle Cell Disease, Boston Medical Center, Boston, Mass (Dr Steinberg); University of Mississippi School of Medicine, Jackson (Drs Steinberg, and Bigelow); Maryland Medical Research Institute, Baltimore (Ms Barton, Dr Terrin); Center for Sickle Cell Disease, Howard University School of Medicine, Washington, DC (Dr Castro); University of Miami School of Medicine, Miami, Fla (Dr Pegelow); Thomas Jefferson University, Philadelphia, Pa (Dr Ballas); Medical College of Georgia, Augusta (Dr Kutlar); University of North Carolina, Chapel Hill (Drs Orringer and Ataga); New York Methodist Hospital, Brooklyn (Dr Bellevue); Hospital for Sick Children, Toronto, Ontario (Dr Olivieri); Sickle Cell Center, Emory University, Atlanta, Ga (Dr Eckman); University of Medicine and Dentistry of New Jersey, Newark (Dr Varma); Roosevelt Medical Center, New York, NY (Dr Ramiez); University of Alabama at Birmingham (Dr Adler); Virginia Commonwealth University, Richmond (Dr Smith); University of Pitts-burgh, Pittsburgh, PA (Dr Carlos); Duke University School of Medicine, Durham, NC (Dr DeCastro); University of Illinois at Chicago (Dr Saithnatharajah); Michael Reese Medical Center, Chicago, Ill (Dr Terrin); Children’s Hospital and Research Center at Oakland, Oakland, Calif (Dr Vichinsky); University of California at San Francisco (Dr Claster); Rainbow Babies and Children’s Hospital, Cleveland, Ohio (Dr Shurin); Brigham and Women’s Hospital, Boston (Dr Bridges); National Heart, Lung, and Blood Institute, Bethesda, Md (Drs Bonds and Waclawiw).

Author Contributions: Study concept and design: Steinberg, Barton, Castro, Ballas, Orringer, Victorin, Shurin, Bridges, Waclawiw, Bonds, Terrin. Acquisition of data: Steinberg, Castro, Ballas, Kutlar, Orringer, Bellevue, Olivier, Varma, Ramirez, Adler, Smith, Carlos, DeCastro, Bigelow, Sauntharajah, Telfer, Claster, Shurin, Bridges, Bonds. Analysis and interpretation of data: Steinberg, Barton, Castro, Ballas, Kutlar, Orringer, Eckman, Victorin, Olivier, Waclawiw, Terrin. Obtained funding: Bonds, Terrin. Administrative, technical, or material support: Castro, Ballas, Kutlar, Ramirez, Adler, Smith, Bigelow, Telfer, Claster, Shurin, Bridges, Bonds, Terrin. Study supervision: Barton, Castro, Ballas, Olivier, Car- los, Vichinsky, Terrin.

Clinical center direction: Telfer, Claster, Shurin, Bridges. Funding/Support: This study was supported by National Heart, Lung, and Blood Institute contract NO1-HB-67129.

Data Safety and Monitoring Board: Lennette Benjam- min, MD; Joel Verter, PhD, Clinton Joiner, MD, PhD; Patricia Adams-Graves, MD; Dorothy Moore, MD.

REFERENCES

209 were from academic books and official reports. The most cited journal was JAMA with 135 citations, followed by the American Journal of Preventive Medicine (102), BMJ (77), and The Lancet (70). Fifty-six journals had articles cited more than 5 times, comprising a total of 1185 citations. Of these 56 journals, 6 (11%) had an impact factor of more than 10.0; 10 (18%) had an impact factor of 5.0 to 10.0; 11 (20%) had an impact factor of 3.0 to 5.0; and 28 (51%) had impact factors of less than 3.0. Of this latter group, 11 (20%) had impact factors of less than 2.0. Only 7 journals (13%) appeared in the top 100 journals ranked by impact factors (2001). The median impact factor of these 56 journals was 2.76. There was a significant correlation between impact factors and times cited in the USPSTF guidelines (Kendall $r = 0.26$, $P = .005$).

**Comment.** We found that the number of citations by the USPSTF guidelines roughly parallels the impact factors for the respective journals. Journals with low impact factors, however, were also cited frequently as providing important evidence. This finding may reflect the fact that journals that focus on preventive services tend to have lower impact factors than do journals in other scientific disciplines.

Some of the possible domains of impact of journal articles that cannot be measured by impact factors are changes in readers' knowledge, practice, clinical outcomes, funding priorities for research, and prompting of further learning. Overreliance on impact factors may undervalue the unique contributions of individual areas of research. In the field of clinical or preventive medicine, in particular, citation analyses on evidence-based practice guidelines may be a more accurate assessment of the contributions of individual journals and researchers. Although we only assessed the area of preventive health services, we suspect that this general conclusion may extend to other areas of scientific inquiry.

Takeo Nakayama, MD, PhD
Department of Medical System Informatics
Tsuguya Fukui, MD, PhD
Department of General Medicine and Clinical Epidemiology

Shunichi Fukuhara, MD, DMsc
Department of Healthcare Research
Kyoto University Graduate School of Medicine
Kyoto, Japan
Kiichiro Tsutani, MD, PhD
Department of Pharmacoconomics
Graduate School of Pharmaceutical Sciences
The University of Tokyo
Tokyo, Japan
Shigeki Yamazaki, PhD
Department of Library and Information Science
Aichi Shukutoku University
Aichi, Japan

**Acknowledgment:** We thank Ms Akiko Yoshida for assistance in editing the manuscript.


**CORRECTIONS**


Incorrect Sentence: In the Original Contribution entitled “Effect of Hydroxyurea on Mortality and Morbidity in Adult Sickle Cell Anemia: Risks and Benefits Up to 9 Years of Treatment” published in the April 2, 2003, issue of THE JOURNAL (2003;289:1645-1651), there was an incorrect sentence on pages 1646 and 1647. The sentence that read “Ninety-six (32%) patients never received hydroxyurea; 48 (16%) received hydroxyurea for less than one year and 156 (52%) received hydroxyurea for 1 or more years” should have read “Thirty-two percent of patients randomly assigned to placebo never received hydroxyurea; 16% received hydroxyurea for less than 1 year and 52% received hydroxyurea for one or more years.”