Effectiveness of Integrated Palliative and Oncology Care for Patients With Acute Myeloid Leukemia: A Randomized Clinical Trial

Areej El-Jawahri, MD; Thomas W. LeBlanc, MD; Alison Kavanaugh, NP; Jason A. Webb, MD; Vicki A. Jackson, MD; Toby C. Campbell, MD; Nina O’Connor, MD; Selina M. Lugter, MD; Ellin Gafford, MD; Jillian Gustin, MD; Bhavana Bhatnagar, DO; Alison R. Walker, MD; Amir T. Fathi, MD; Andrew M. Brunner, MD; Gabriela S. Hobbs, MD; Showly Nicholson, BS; Debra Davis, RN, BSN; Hilena Addis, BS; Dagny Vaughn, BA; Nora Horick, MS; Joseph A Greer, PhD; Jennifer S. Temel, MD

IMPORTANCE Patients with acute myeloid leukemia (AML) receiving intensive chemotherapy experience substantial decline in their quality of life (QOL) and mood during their hospitalization for induction chemotherapy and often receive aggressive care at the end of life (EOL). However, the role of specialty palliative care for improving the QOL and care for this population is currently unknown.

OBJECTIVE To assess the effect of integrated palliative and oncology care (IPC) on patient-reported and EOL outcomes in patients with AML.

DESIGN, SETTING, AND PARTICIPANTS We conducted a multisite randomized clinical trial of IPC (n = 86) vs usual care (UC) (n = 74) for patients with AML undergoing intensive chemotherapy. Data were collected from January 2017 through July 2019 at 4 tertiary care academic hospitals in the United States.

INTERVENTIONS Patients assigned to IPC were seen by palliative care clinicians at least twice per week during their initial and subsequent hospitalizations.

MAIN OUTCOMES AND MEASURES Patients completed the 44-item Functional Assessment of Cancer Therapy–Leukemia scale (score range, 0-176) to assess QOL; the 14-item Hospital Anxiety and Depression Scale (HADS), with subscales assessing symptoms of anxiety and depression (score range, 0-21); and the PTSD Checklist–Civilian version to assess posttraumatic stress disorder (PTSD) symptoms (score range, 17-85) at baseline and weeks 2, 4, 12, and 24. The primary end point was QOL at week 2. We used analysis of covariance adjusting and mixed linear effect models to evaluate patient-reported outcomes. We used Fisher exact test to compare patient-reported discussion of EOL care preferences and receipt of chemotherapy in the last 30 days of life.

RESULTS Of 235 eligible patients, 160 (68.1%) were enrolled; of the 160 participants, the median (range) age was 64.4 (19.7-80.1) years, and 64 (40.0%) were women. Compared with those receiving UC, IPC participants reported better QOL (adjusted mean score, 107.59 vs 116.45; P = .04), and lower depression (adjusted mean score, 7.20 vs 5.68; P = .02), anxiety (adjusted mean score, 5.94 vs 4.53; P = .02), and PTSD symptoms (adjusted mean score, 31.69 vs 27.79; P = .01) at week 2. Intervention effects were sustained to week 24 for QOL (β, 2.35; 95% CI, 0.02-4.68; P = .048), depression (β, −0.42; 95% CI, −0.82 to −0.02; P = .04), anxiety (β, −0.38; 95% CI, −0.75 to −0.01; P = .04), and PTSD symptoms (β, −1.43; 95% CI, −2.34 to −0.54; P = .002). Among patients who died, those receiving IPC were more likely than those receiving UC to report discussing EOL care preferences (21 of 28 [75.0%] vs 12 of 30 [40.0%]; P = .01) and less likely to receive chemotherapy near EOL (15 of 43 [34.9%] vs 27 of 41 [65.9%]; P = .01).

CONCLUSIONS AND RELEVANCE In this randomized clinical trial of patients with AML, IPC led to substantial improvements in QOL, psychological distress, and EOL care. Palliative care should be considered a new standard of care for patients with AML.

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Patients with acute myeloid leukemia (AML) receiving intensive induction chemotherapy face an abrupt onset of a life-threatening illness that necessitates urgent initiation of treatment and a prolonged hospitalization for recovery.\textsuperscript{1-5} During this hospitalization, patients experience marked physical symptoms due to effects of intensive chemotherapy, which negatively effects their quality of life (QOL).\textsuperscript{6-12} The majority of patients also experience psychological distress as they struggle with uncertainty regarding their prognosis, the isolation they experience during the hospital stay, and the loss of their independence.\textsuperscript{6-19} In addition to their physical and psychosocial symptom burden, there is a critical need to optimize end-of-life (EOL) care for patients with AML.\textsuperscript{20-24} These patients rarely discuss their EOL care preferences, are often hospitalized, and receive chemotherapy in the last weeks of life.\textsuperscript{18,21,25,26} Yet, interventions to improve QOL, reduce psychological distress, and optimize EOL care for this population are lacking.

Specialty palliative care has been shown to improve QOL, reduce symptom burden and psychological distress, and enhance EOL outcomes for patients with advanced solid tumors.\textsuperscript{27-31} Additionally, palliative care integrated with transplant care has been shown to improve QOL and reduce psychological distress for patients with hematologic cancers undergoing stem cell transplantation.\textsuperscript{32,33} However, oncologists rarely consult palliative care for patients with AML, in part because of the lack of evidence for the role of early palliative care in this population.\textsuperscript{21,22,34,35} Clinical trials are needed to determine if involvement of palliative care can improve the experience and outcomes of patients with AML.

We conducted a multisite, nonblinded randomized clinical trial to assess the effect of integrated palliative and oncology care (IPC) vs usual care (UC) on QOL, mood, symptom burden, and EOL outcomes for hospitalized patients with AML receiving intensive chemotherapy. We hypothesized that patients receiving IPC would have (1) better QOL, (2) lower psychological distress, (3) reduced symptom burden, (4) higher rates of discussing their EOL care preferences with their clinicians, and (5) lower rates of hospitalization and chemotherapy administration near the EOL.

**Methods**

**Participants**

Hospitalized patients 18 years and older with high-risk AML receiving intensive chemotherapy were eligible to participate. We defined patients with high-risk AML as (1) newly diagnosed patients 60 years and older with an antecedent hematologic disorder or therapy-related disease, or (2) patients with relapsed or primary refractory AML. We considered intensive chemotherapy as a combination of anthracycline and cytarabine (ie, the 7 + 3 regimen) or a modification of this regimen on a clinical trial with additional drug(s) added or other similar intensive chemotherapy regimens requiring a 3- to 6-week hospitalization. We excluded patients with a diagnosis of acute promyelocytic leukemia and those receiving non-intensive chemotherapy. We also excluded patients already receiving palliative care and those with major psychiatric or comorbid conditions that would prohibit their adherence to study procedures, as determined by the treating oncologist.

**Integrated Palliative and Oncology Care Intervention**

Patients randomized to IPC met with an inpatient palliative care physician, advance practice nurse, or physician assistant within 72 hours of randomization. The palliative care clinician conducted at least 2 visits per week throughout the patient’s hospitalization for intensive chemotherapy and all subsequent hospitalizations up to 1 year after randomization. Patients and the palliative care clinician were permitted to initiate additional visits during hospitalizations as needed. Palliative care clinicians did not see patients in the outpatient setting.

We developed the IPC intervention based on prior work developing and evaluating the effect of palliative care for patients with solid tumors, and also those with hematologic cancers undergoing stem cell transplantation.\textsuperscript{27,28,32,33} Palliative care clinicians conducted at least 2 visits per week throughout the patient’s hospitalization for intensive chemotherapy and all subsequent hospitalizations up to 1 year after randomization. Patients and the palliative care clinician were permitted to initiate additional visits during hospitalizations as needed. Palliative care clinicians did not see patients in the outpatient setting.
Care clinicians initially focused on establishing rapport, assessing palliative care needs, and developing a relationship with the patient. Throughout hospitalization, clinicians addressed patients’ symptoms, assessed their illness understanding, ascertained their goals and expectations, and assisted with their treatment decision-making. Palliative care clinicians documented the elements of care that they addressed after each visit by using a structured questionnaire in the Research Electronic Data Capture (REDCap) system.

**Usual Care**
Patients assigned to UC received supportive care measures as per their oncology team. They were permitted to receive palliative care at their request or at the request of their oncologist.

**Study Measures**

**Participant-Reported Measures**
We used the 44-item Functional Assessment of Cancer Therapy-Leukemia scale that includes 5 subscales assessing physical, functional, emotional, social well-being, and leukemia-specific concerns during the past week (score range, 0-176), with higher scores indicating better QOL. 36 We measured patients’ anxiety and depression with the 14-item Hospital Anxiety and Depression Scale (HADS). The HADS consists of 2 subscales assessing symptoms of anxiety and depression, with subscale scores ranging from 0 (no distress) to 21 (maximum distress) and cut-off scores of more than 7 indicating clinically significant symptoms. 37 We also assessed depression with the 9-item Patient Health Questionnaire (PHQ-9), a measure that detects symptoms of major depressive disorder, according to the criteria of the Diagnostic and Statistical Manual of Mental Disorders, and can be evaluated continuously with higher scores indicating worse mood. 38 We used the revised 10-item Edmonton Symptom Assessment Scale, which uses a 0-10 scale (score range, 0-100), with higher scores indicating greater symptom burden. 39 We used the 17-item PTSD Checklist-Civilian version to evaluate the severity of posttraumatic stress disorder (PTSD) symptoms (score range, 17-85), with higher scores indicating worse PTSD symptoms. 40

We used 1 item to assess patient-reported discussion of EOL care preferences with their clinicians, as per prior studies. 31 Specifically, patients were asked, “Have you and your doctors discussed any particular wishes you have about the care you want to receive if you were dying?” Response items were yes or no. Although patients completed this measure at all study time points, we used the assessment prior to death or at 6-month follow-up, as defined in the study protocol (Supplement 1).

**EOL Outcomes**
The research assistant collected data on hospitalization, chemotherapy administration based on chemotherapy administration flow sheets, and hospice referrals (if applicable) from patient electronic medical records (EMRs). The research assistant obtained date of death from the patient EMRs or from obituaries. All EOL outcomes were obtained by January 2020, at a minimum of 6-month follow-up for all study participants.

**Statistical Analysis**
We performed statistical analyses using Stata, version 9.3 (StataCorp). We summarized participants’ baseline characteristics between randomized groups, using frequency and percentage for categorical variables and median (SD) range for continuous variables. A 2-sided P value less than .05 was considered statistically significant.

The primary end point of the study was a comparison of QOL at week 2 between study groups using analysis of covariance and controlling for baseline criterion score. We chose the second week of hospitalization as the primary end point because it is the most symptomatic phase of the intensive chemotherapy hospitalization. 10,42 We also compared symptom burden using the Edmonton Symptom Assessment Scale, depression symptoms using PHQ-9 and the depression subscale of HADS, anxiety symptoms using the anxiety subscale of HADS, and PTSD symptoms using the PTSD Checklist-Civilian at week 2 between the study groups using analysis of covariance and controlling for baseline criterion score. We also dichotomized the HADS depression and anxiety subscale scores as described above to compare frequencies of depression and anxiety symptoms between the study groups at week 2 using Fisher exact test.

We then used mixed linear effect models using maximum likelihood to account for missing data to examine the effect of IPC on patient-reported outcomes longitudinally across all time points (baseline, week 2, week 4, week 12, and week 24). For all these analyses, we report the beta estimated coefficient (β). A positive β coefficient indicates a positive association between the intervention and the outcome of interest. A negative β coefficient indicates a negative association between the intervention and the outcome of interest.

We compared rates of patient-reported EOL discussions, hospitalizations in the last week of life, chemotherapy administration in the last 30 days of life, and hospice use between the 2 study groups by using Fisher exact test. We used Poisson regression to compare hospice length of stay between the study groups.

We powered this study based on prior experience integrating palliative care for patients undergoing stem cell transplantation. With a sample size of 160 patients, we ensured greater than 90% power to detect a 6.9-point difference in patient-reported QOL at week 2, with a 2-sided .05 significant level and assuming 10% missing data at week 2. To address the issue of multiple tests, we used the false discovery rate (FDR) control method. For secondary outcomes, we selected an FDR of 15%, which denotes the acceptable percentage of results that potentially represent false positives. 43
randomization, 86 patients were assigned to IPC, and 74 patients were assigned to UC. Of the 160 enrolled patients, 138 were White (86.2%), and the median (range) age was 64.4 (19.7-80.1) years (eTable 1 in Supplement 2). Overall, 109 (68.1%) patients had newly diagnosed AML. There were no meaningful differences in clinical characteristics between the 2 study groups. At weeks 2, 4, 12, and 24, data were missing for 13 (8.1%), 23 (14.4%), 23 (14.4%), and 55 (34.4%) patients, respectively.

**Palliative Care Visits**

Patients assigned to IPC had a mean (range) of 2.2 (2-5) visits per week during their hospitalization for intensive chemotherapy. Only 6 patients assigned to UC received a palliative care consultation during their initial hospitalization, and 24 of 74 (32.4%) patients had a palliative care consultation during subsequent hospitalizations. During hospitalization for intensive chemotherapy, the palliative care clinicians most commonly reported establishing rapport (64.9%), addressing symptoms (64.7%), and coping (63.6%) with patients receiving the intervention (eFigure 1 in Supplement 2); eFigure 2 in Supplement 2 depicts topics covered by palliative care clinicians during subsequent hospitalizations.

**Patient-Reported Outcomes at Week 2**

Patients assigned to IPC, when compared with patients assigned to UC, reported better QOL (adjusted mean score, 116.45 vs 107.59; \( P = .04 \)), lower depression (HADS depression subscale: adjusted mean score, 5.68 vs 7.20; \( P = .02 \)); and PHQ-9: adjusted mean score, 6.34 vs 8.00; \( P = .04 \)), anxiety (adjusted mean score, 4.53 vs 5.94; \( P = .02 \)), and PTSD symptoms (adjusted mean score, 27.79 vs 31.69; \( P = .01 \)) at week 2 (Table). There were no differences in symptom burden between the 2 groups. The findings remain statistically significant when using FDR to correct for multiple testing (eTable 2 in Supplement 2). At week 2, patients assigned to IPC, when compared with those receiving UC, also reported lower rates of clinically significant symptoms of depression (22 of 78 [28.2%] vs 31 of 69 [44.9%]; \( P = .04 \)) and anxiety (17 of 78 [21.8%] vs 26 of 96 [37.7%]; \( P = .046 \)).

**Longitudinal Assessment of Patient-Reported Outcomes**

Using mixed linear effects models with maximum likelihood to impute missing data, when compared to patients receiving UC, patients assigned to IPC reported better QOL (\( \beta = 2.35; 95\% \ CI, 0.02-4.68; P = .048 \)), lower depression (HADS depression subscale: \( \beta = -0.42; 95\% \ CI, -0.82 to -0.02; P = .04 \)), anxiety (\( \beta = -0.38; 95\% \ CI, -0.75 to -0.01; P = .04 \)), and PTSD symptoms (\( \beta = -1.43; 95\% \ CI, -2.34 to -0.54; P = .002 \)) longitudinally (Figure 2). There were no differences in symptom burden or PHQ-9 scores longitudinally. The findings remained statistically significant when using FDR to correct for multiple testing (eTable 2 in Supplement 2).

**End-of-Life Outcomes**

Among patients who died (n = 87; 44 of 73 in the UC group and 43 of 84 in the IPC group), those receiving IPC vs UC were more likely to report discussing their EOL care preferences with their clinicians (21 of 28 [75.0%] vs 12 of 30 [40.0%]; \( P = .01 \)) and less likely to receive chemotherapy in the last 30 days of life (15 of 43 [34.9%] vs 27 of 41 [65.9%]; \( P = .01 \)). Among those receiving IPC vs UC, there was no difference in hospice use (15 of 41 [34.9%] vs 15 of 42 [36.6%]; \( P = .999 \)), hospice length of stay (\( \beta = -0.80; 95\% \ CI, -1.85 to 0.25; P = .14 \)), and hospitalization in the last week of life (35 of 42 [83.3%] vs 32 of 43 [74.4%]; \( P = .43 \)). The findings remain statistically significant when using FDR to correct for multiple testing (eTable 2 in Supplement 2).
Discussion

Results of this multisite randomized clinical trial demonstrate that IPC improves QOL, depression and anxiety symptoms, and posttraumatic stress symptoms for patients with AML receiving intensive chemotherapy compared with usual care. The intervention led to clinically meaningful and sustained improvements in QOL and psychological distress for 6 months after initiating chemotherapy in this population at

Table. Effect of Integrated Palliative and Oncology Care on Patient-Reported Outcomes at Week 2

<table>
<thead>
<tr>
<th>Measure (scale)</th>
<th>Sample size</th>
<th>Group assignment</th>
<th>Adjusted mean score (95% CI)</th>
<th>Standardized mean differencea</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Quality of life (FACT-Leukemia)</td>
<td>139</td>
<td>Usual care</td>
<td>107.59 (101.45-113.74)</td>
<td>0.30</td>
<td>.04</td>
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<tr>
<td></td>
<td></td>
<td>Integrated palliative and oncology care</td>
<td>116.45 (110.69-122.21)</td>
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<td></td>
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<tr>
<td>Anxiety symptoms (HADS)</td>
<td>147</td>
<td>Usual care</td>
<td>5.94 (5.10-6.79)</td>
<td>0.31</td>
<td>.02</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Integrated palliative and oncology care</td>
<td>4.53 (3.74-5.33)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Depression symptoms (HADS)</td>
<td>147</td>
<td>Usual care</td>
<td>7.20 (6.26-8.14)</td>
<td>0.34</td>
<td>.02</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Integrated palliative and oncology care</td>
<td>5.68 (4.80-6.56)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Depression syndrome (PHQ-9)</td>
<td>144</td>
<td>Usual care</td>
<td>8.00 (6.83-9.17)</td>
<td>0.31</td>
<td>.04</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Integrated palliative and oncology care</td>
<td>6.34 (5.23-7.44)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Symptom burden (ESAS)</td>
<td>146</td>
<td>Usual care</td>
<td>32.82 (28.58-37.06)</td>
<td>0.23</td>
<td>.12</td>
</tr>
<tr>
<td>PTSD symptoms (PTSD Checklist–Civilian)</td>
<td>146</td>
<td>Usual care</td>
<td>31.69 (29.56-33.82)</td>
<td>0.30</td>
<td>.01</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Integrated palliative and oncology care</td>
<td>27.79 (25.78-29.80)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: ESAS, Edmonton Symptom Assessment Scale; FACT–Leukemia, Functional Assessment of Cancer Therapy–Leukemia; HADS, Hospital Anxiety and Depression Scale; PHQ-9, 9-item Patient-Health Questionnaire; PTSD, posttraumatic stress disorder.

When compared with patients assigned to usual care, patients assigned to integrated palliative and oncology care reported better quality of life (β, 2.35; 95% CI, 0.02-4.68; P = .048) (A), lower anxiety (β, −0.38; 95% CI, −0.75 to −0.01; P = .04) (B), lower depression (β, −0.42; 95% CI, −0.82 to −0.02; P = .04) (C), and fewer posttraumatic stress disorder (PTSD) symptoms (β, −1.43; 95% CI, −2.34 to −0.54; P = .002) (D).
high risk for long-term QOL impairments and psychological morbidity. Although many oncologists question palliative care clinicians’ ability to meet the specialized needs of patients with AML, the present findings provide compelling evidence to the contrary.22,26,34,35,44-46 Although prior randomized trials of IPC care models have traditionally excluded patients with hematologic cancers,27-29,31 this study establishes the role of IPC care for improving the QOL and care in patients with AML. While we cannot generalize these findings to all patients with hematologic cancers, the salient benefits of early palliative care in patients hospitalized with AML are consistent to what we see in patients with hematologic cancers undergoing stem cell transplantation.32,33 In contrast with the prior randomized trial of integrated palliative care for patients undergoing stem cell transplant,32,33 this study did not demonstrate a statistically significant difference in symptom burden between the 2 groups. The extent of symptom burden for patients receiving induction chemotherapy in this trial was high and comparable with those seen in prior studies of this population and those undergoing myeloablative stem cell transplantation.11,47-49 Nonetheless, findings of this trial provide compelling evidence to support palliative care integration into routine clinical care for patients with high-risk hematologic cancers, especially those enduring prolonged hospitalizations.

Psychological distress during intensive hospitalizations for patients with hematologic cancers is associated with long-term sequelae, including psychiatric morbidity, medical complications, and even mortality.18,19,50-54 Involvement of palliative care clinicians in the care of patients with AML led to notable improvement in all psychological outcomes, including depression, anxiety, and posttraumatic stress symptoms. Strikingly, these improvements were sustained up to 6 months after initiating therapy in this population. Given the growing literature on traumatic stress and PTSD in patients with AML, literature on traumatic stress and PTSD in patients with acute myeloid leukemia (AML), the present findings provide compelling evidence to the contrary.22,26,34,35,44-46 Although prior randomized trials of IPC care models have traditionally excluded patients with hematologic cancers,27-29,31 this study establishes the role of IPC care for improving the QOL and care in patients with AML. While we cannot generalize these findings to all patients with hematologic cancers, the salient benefits of early palliative care in patients hospitalized with AML are consistent to what we see in patients with hematologic cancers undergoing stem cell transplantation.32,33 In contrast with the prior randomized trial of integrated palliative care for patients undergoing stem cell transplant,32,33 this study did not demonstrate a statistically significant difference in symptom burden between the 2 groups. The extent of symptom burden for patients receiving induction chemotherapy in this trial was high and comparable with those seen in prior studies of this population and those undergoing myeloablative stem cell transplantation.11,47-49 Nonetheless, findings of this trial provide compelling evidence to support palliative care integration into routine clinical care for patients with high-risk hematologic cancers, especially those enduring prolonged hospitalizations.

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Patients receiving the palliative care intervention also experienced improvements in critical EOL outcomes. Specifically, patients receiving the intervention were more likely to discuss their EOL care preferences with their clinicians and less likely to use chemotherapy in the last month of life. More than half of the study cohort died during the study period, with the majority of patients receiving chemotherapy in the last 30 days of life and hospitalized in the last week of life, which highlights the poor prognosis and intensity of EOL care that has been described in this population.18,21,25,26,58-61 While there were no differences in hospitalizations near the EOL or hospice use between study groups, this likely reflects the fact that hospice services are not well equipped to meet the EOL needs of this population, including palliative transfusions.20,22,26,59-62 In fact, the American Society of Hematology has released a statement recommending that hospice agencies and payers work collaboratively to ensure the availability of palliative transfusions to optimize EOL care for patients with hematologic cancers.63 Future studies examining innovative EOL care delivery models are needed to minimize the need for hospitalizations in this population.

Limitations
This study has several limitations. First, it was performed only at tertiary care academic centers. While this is the predominant setting where intensive chemotherapy is given, these findings may not be generalizable to other care settings. Second, the sample lacked racial and ethnic diversity, and thus we are unable to assess the effect of these important factors on study outcomes. Third, study staff, patients, and clinicians could not be blinded to the intervention, which may have introduced bias. Although prior studies have suggested low risk of bias with self-reported assessments and objective EOL metrics obtained from patient EMRs,64-66 this is nonetheless an important study limitation. Fourth, the intervention did not include the full interdisciplinary palliative care team, which may have led to more profound benefits for this population. Fifth, similar to prior palliative care studies,27-33 we did not use an attention control group to adjust for the potential benefits of the time palliative care clinicians spent with patients. Finally, the involvement of palliative care in the care of patients randomized to the intervention on the same leukemia hospital floor may have altered clinicians and nursing behaviors in the control group, which may have diluted the findings. Additionally, a substantial proportion of patients in the control group received palliative care during their illness course, which may have also diluted the findings.

Conclusions
The American Society of Clinical Oncology recommends concurrent palliative care from the time of diagnosis for all patients with metastatic cancer and/or high symptom burden.67,68 However, the role of palliative care in the care of patients with AML has remained uncertain given the lack of evidence of benefit for this population. Results of this randomized clinical trial demonstrate that early IPC for hospitalized patients with AML receiving intensive chemotherapy can substantially enhance their QOL and reduce their depression, anxiety, and posttraumatic stress symptoms during hospitalization for intensive chemotherapy and up to 6 months after diagnosis. Patients receiving the IPC model were also more likely to experience improvements in their EOL care. Importantly, induction chemotherapy is offered mostly at large academic hospitals with access to inpatient palliative care services, which allows for potential implementation and dissemination of this care model for patients with AML. As these patients spend the majority of their time in the hospital and clinical settings, there are numerous opportunities to engage palliative care clinicians early and longitudinally in their care. Thus, early palliative care at the time of diagnosis for patients with AML should become standard of care to improve the QOL and care for this population.
Research

Effectiveness of Integrated Palliative and Oncology Care for Patients With Acute Myeloid Leukemia

ARTICLE INFORMATION

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