Fludarabine + prednisone ± α-interferon followed or not by α-interferon maintenance therapy for previously untreated patients with chronic lymphocytic leukemia: long term results of a randomized study

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Background and Objectives: Fludarabine is an effective therapy for patients with chronic lymphocytic leukemia (CLL) and interferon-α (IFN-α) has been reported to have anti-leukemic activity in CLL patients. A randomized study was designed to evaluate whether the addition of IFN-α to a first-line treatment with fludarabine and prednisone could increase the response rate in patients with advanced CLL and whether IFN-α given as maintenance therapy could improve the duration of response.

Design and Methods. One hundred and thirty-three patients were randomized to receive fludarabine (25 mg/m² i.v. days 9-13) and prednisone (20 mg/m², days 1, 3, 5, 7 and 14 and 40 mg/m², days 9-13) (arm A: 66 patients) or in addition to the same schedule, IFN-α (2 MU/ sc, days 1, 3, 5, 7, 9, 11, 13 and 15) (arm B: 67 patients). Seventy-eight patients responsive to therapy entered the post-remission phase of the study in which 41 patients were randomized to receive IFN-α (3 MU three times a week) and 37 to clinical observation.

Results. A similar response rate (complete responses + partial responses + partial remissions) was observed in the 2 arms: 86% for arm A and 84% for arm B (p = 0.4). A longer duration of the response observed in patients who achieved a complete response (p = 0.001) and in patients who received maintenance therapy with IFN-α (p < 0.05). However, the quality of response was the only significant and independent factor influencing response duration (p < 0.01). No benefits in terms of infection-related mortality and morbidity could be ascribed to IFN-α administration.

Interpretation and Conclusions. In previously untreated CLL patients with advanced disease a high response rate is obtained from first-line fludarabine and prednisone and no benefit is derived from the addition of IFN-α to this regimen. The achievement of a good quality response to therapy was the only independent predictor of a prolonged response.

Key words: chronic lymphocytic leukemia, treatment, fludarabine, α-interferon.

Haematologica 2003; 88:1348-1357
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During the last 10 years three prospective randomized studies have demonstrated that fludarabine therapy is superior to CAP (cyclophosphamide, doxorubicin, and prednisone),1 CHOP (cyclophosphamide, vincristine, doxorubicin, and prednisone)2 and chlorambucil3 as first-line treatment of patients with chronic lymphocytic leukemia (CLL). However, most patients relapse after 2-4 years suggesting that fludarabine has a very effective debulking activity, but that it is not curative: residual disease represents the main cause of relapse.

Interferon-α (IFN-α) is a biological agent with multiple properties including inhibition of cell proliferation and modulation of cellular immunity.4-5 IFN-α has been shown to have an anti-tumor effect in different hematologic malignancies, in particular multiple myeloma, non-Hodgkin’s lymphoma and hairy cell leukemia.6-12 During the late eighties, several clinical trials evaluated the anti-tumor activity of IFN-α, given at variable doses and for different durations, as a single agent in CLL patients with different clinical pictures. While only limited activity was recorded in patients previously treated and with advanced disease,13-18 a higher response rate was observed in untreated patients with early disease.13-22 Furthermore, some authors reported a therapeutic benefit of IFN-α given as post-remission therapy.23,24

A synergistic effect of IFN-α with several cytotoxic drugs has been documented in vitro25 and favorable clinical results in patients with low grade lymphoma have been described.26 Positive clinical results were reported by Mandelli et al.27 and by Molica et al.28 in small series of CLL patients with advanced disease treated with a regimen including chlorambucil, prednisone and IFN-α.

On the basis of the anti-leukemic activity shown by IFN-α given alone or in combination with cytotoxic drugs and considering its potential immuno-modulatory properties, a multicenter study was designed to evaluate the therapeutic effect of IFN-α in previously untreated CLL patients with advanced disease. The study was characterized by two randomized phases. In the first phase, the activity and toxicity of two induction schedules, fludarabine and prednisone with or without IFN-α, were evaluated. The primary objective of this first part of the study, which included 134 CLL patients, was to evaluate whether the addition of IFN-α could increase the response rate. In the second phase, patients who achieved a response to induction therapy were randomized to receive IFN-α as maintenance therapy or no treatment. The primary objective of this second part of
the study, which included 78 CLL patients, was to evaluate whether the administration of IFN-α as maintenance therapy could improve the response duration. Herein, we report, the results of this study.

**Design and Methods**

**Patients**

One hundred and thirty-four patients with untreated progressive or advanced CLL were prospectively enrolled into a randomized clinical trial between March 1993 and December 1998. Patients were recruited from three Italian hematologic centers: Dipartimento di Biotecnologie Cellulari ed Ematologia, University “La Sapienza” of Rome (74 patients), Istituto di Ematologia ed Oncologia “L & A. Seragcoli”, University of Bologna (32 patients), and Istituto di Ematologia, University of Udine (28 patients).

**Pre-study evaluation and eligibility criteria**

All patients fulfilled the National Cancer Institute-Sponsored Working-Group diagnostic criteria for CLL. At baseline, peripheral blood lymphocytes were characterized by immunophenotyping and morphology as previously described. The stage of the disease was assessed according to Rai’s classification. Pre-treatment work-up included a medical history, physical examination, complete peripheral blood (PB) cell count with differential, bone marrow (BM) aspiration and biopsy, Ig quantification, liver and renal function tests and radiographic examination (computerized tomography scans, ultrasounds).

Eligibility criteria included: age 65 years or less, no prior treatment, advanced stage (III-IV) or intermediate stage (I-II) with one or more clinical signs of active disease. Exclusion criteria included: prior treatment; autoimmune cytopenia; history of other malignancies within 2 years prior to study entry (except for adequately treated carcinoma in situ of the cervix; basal or squamous cell skin cancer); active infection requiring systemic therapy, human immunodeficiency virus infection, hepatitis B or C; history of uncontrolled hypertension, severe cardiac, pulmonary, or neurological disease; uncontrolled metabolic disorder; any co-existing medical or psychological condition that would preclude participation in the study or compromise ability to give informed consent.

The protocol was approved by the local ethics committee. All patients gave written informed consent before enrollment into the study which was carried out in accordance with the Good Clinical Practice precepts.

**Treatment plan**

Eligible patients were centrally randomized to one of the two induction treatments (arm A or arm B).

Randomization was balanced for each participating center and stratified according to the following factors: institution, stage and bone marrow histology. Arm A: fludarabine (Fludara® kindly given by Schering SpA) associated with prednisone. Fludarabine: 25 mg/m² i.v. for 5 consecutive days, on days 9 to 13; Prednisone: 20 mg/m² orally on days 1, 3, 5, 7 and 14. Prednisone: 40 mg/m² orally from day 9 to day 13.

Arm B: fludarabine associated with prednisone as in arm A with the addition of IFN-α as follows: IFN-α: (lymphoblastoid IFN-α; Wellferon® kindly given by Glaxo-Wellcome) 2 MUI on days 1, 3, 5, 7, 9, 11, 13 and 15.

Both regimens were administered every 4 weeks for a total number of 6 courses. Patients achieving a complete or partial response (CR or PR) according to NCI criteria were considered for the second randomized part of the study, the post-remission phase. Patients were randomized to two post-remission approaches (arm C or arm D).

Arm C: IFN-α: 3 MUI three times a week until disease progression

Arm D: no therapy.

Randomization was stratified according to 3 factors: institution, prior induction therapy and quality of response. All study treatments were carried out on an outpatient basis and IFN-α was self-administered subcutaneously.

**Dose modifications**

In the presence of severe (III-IV grade according to WHO criteria) hematologic toxicity, the start of the subsequent course was delayed. In the case of severe cytopenias persisting for more than 2 weeks, treatment was discontinued, while, in the presence of moderate cytopenias, fludarabine and IFN-α doses were reduced by 50%. Treatment was withheld in the presence of grade III-IV non-hematologic WHO toxicity or major infection until the patient had recovered from toxicity or infection. In the case of recovery after more than 2 weeks or no recovery, patients were withdrawn from the study.

**Supportive care**

During fludarabine treatment and for at least 1 year after therapy discontinuation, trimethoprim-sulphamethoxazole was given three times a week for Pneumocystis carinii prophylaxis. No granulocyte colony-stimulating factor (G-CSF) administration was planned in the presence of neutropenia. Patients requiring blood transfusions were given irradiated products. To prevent flu-like symptoms, paracetamol was administered 30 minutes prior to IFN-α.
Response and toxicity evaluation

Response to induction therapy was assessed according to NCI criteria.\(^2^9\) Response duration was measured from the time of achievement of the response to the time of disease progression. Disease progression was based on the presence on two monthly consecutive evaluations of one or more of the following disease-related signs: ≥50% increase in the absolute number of circulating lymphocytes (minimum number: at least 5000/µL); ≥50% increase of the size of spleen, liver and lymph-nodes (minimum diameter: at least 2 cm). Disease transformation was considered in the presence of a histologic diagnosis of lymphoma and in the case of an increase of the prolymphocyte rate ≥55% (prolymphocytoid transformation). Autoimmune hemolytic anemia was considered to have developed when there were clinical signs of hemolysis associated with a positive Coombs' test. Toxicity was evaluated according to WHO criteria. Toxicity was separately recorded and evaluated in the two phases of the study, during the induction therapy and the post-remission phase, from response to the start of a second line therapy.

**Statistical analysis**

Statistical analysis included an evaluation of the response rate, of the actuarial time to progression probability and of the actuarial survival probability. The corrected \(\chi^2\) test was applied to compare groups. Survival curves were calculated according to Kaplan and Meier\(^3^2\) from the time of first randomization and from the time of second randomization to death, and compared with the log-rank test.\(^3^3\) Response duration was calculated from the time of response to the time of disease progression or death.

The prognostic significance of the following parameters: gender (male vs female), age (≤55 vs >55 years), time from CLL diagnosis to treatment (≤12 vs >12 months), stage according to the Rai classification (I+II vs III+IV), peripheral blood lymphocyte count (≤60 vs >60×10^9/L), LDH value (normal vs elevated), rate of peripheral blood lymphocyte reduction after the second course of induction therapy (≤25 vs >25%), BM histology (non-diffuse vs diffuse), induction therapy regimen (fludarabine + prednisone vs fludarabine + prednisone + IFN-\(\alpha\): arm A vs arm B), on the probability of achieving a response was analyzed. Furthermore, the prognostic significance on response duration and survival of these same parameters, quality of response (CR vs PR) and type of post-remission approach (IFN-\(\alpha\) vs no treatment) was evaluated.

In order to evaluate the relative significance of prognostic factors emerging as such from the univariate analysis, the multiple regression model of Cox was applied.\(^3^4\)

**Results**

**Clinical features of patients**

The median age of the 134 CLL patients entered into the study was 54 years (range: 34-65 years). The median duration of CLL before the start of treatment was 12.6 months. More than two-thirds of patients were males and 13% had Rai stage III-IV disease. The median hemoglobin value was 13 g/dL, the median lymphocyte count 67×10^9/L and the median platelet count 157×10^9/L. The patients' characteristics are reported in Table 1.

**Induction therapy**

Sixty-seven patients were randomized to receive fludarabine and prednisone (arm A) and 67 were randomized to receive fludarabine, prednisone and IFN-\(\alpha\) (arm B). The baseline clinical features of the

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### Table 1. Patients' characteristics.

<table>
<thead>
<tr>
<th>No. of patients</th>
<th>FLU+PDN 67 (%)</th>
<th>FLU+PDN +IFN-(\alpha) 67 (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>94 (70)</td>
<td>45 (67)</td>
</tr>
<tr>
<td>Female</td>
<td>40 (30)</td>
<td>22 (33)</td>
</tr>
<tr>
<td>Median age (yrs.)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>54-65</td>
<td>34-65</td>
</tr>
<tr>
<td>Median time from diagnosis (months)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>0-103</td>
<td>0-103</td>
</tr>
<tr>
<td>Rai stage</td>
<td></td>
<td></td>
</tr>
<tr>
<td>I+II</td>
<td>117 (87)</td>
<td>59 (88)</td>
</tr>
<tr>
<td>III+IV</td>
<td>19 (13)</td>
<td>8 (12)</td>
</tr>
<tr>
<td>Median hemoglobin (g/dL)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>7.1-17.4</td>
<td>7.1-17.4</td>
</tr>
<tr>
<td>Median lymphocytes (×10^9/L)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>2-315</td>
<td>9-167</td>
</tr>
<tr>
<td>Median platelets (×10^9/L)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>70-410</td>
<td>72-318</td>
</tr>
<tr>
<td>Bone marrow histology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diffuse</td>
<td>58 (43)</td>
<td>30 (45)</td>
</tr>
<tr>
<td>Non-diffuse</td>
<td>76 (57)</td>
<td>37 (55)</td>
</tr>
<tr>
<td>LDH</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>94 (70)</td>
<td>49 (73)</td>
</tr>
<tr>
<td>Elevated</td>
<td>40 (30)</td>
<td>22 (33)</td>
</tr>
</tbody>
</table>

NS: not significant.
Table 2. Response to therapy by induction therapy arm.

<table>
<thead>
<tr>
<th></th>
<th>No. of patients</th>
<th>FLU+PDN</th>
<th>FLU+PDN plus IFN-α</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall responses</td>
<td>133 (85)</td>
<td>57 (86)</td>
<td>56 (84)</td>
<td>NS</td>
</tr>
<tr>
<td>CR</td>
<td>50 (38)</td>
<td>30 (45)</td>
<td>20 (30)</td>
<td>NS</td>
</tr>
<tr>
<td>PR</td>
<td>63 (47)</td>
<td>27 (41)</td>
<td>36 (54)</td>
<td></td>
</tr>
</tbody>
</table>

NS: not significant.

Two groups of patients did not differ except for the higher median age ($p < 0.01$) of patients randomized to receive no IFN-α (Table 1).

Response to induction therapy. One patient in whom the immunologic characteristics did not fulfill the criteria for a CLL diagnosis was retrospectively excluded from the study. No protocol deviations were recorded in the induction phase. Thus, response was assessed in 133 patients. The overall response rate (CR+PR) was 85% (113 patients) with no statistically significant differences between the two arms (arm A, fludarabine, prednisone: 86% vs arm B, fludarabine, prednisone and IFN-α: 84%; $p = 0.4$).

A higher, though not significantly so, CR rate was observed in patients treated with fludarabine and prednisone compared to patients treated with these two drugs and IFN-α (45% vs 30%; $p = 0.08$) (Table 2).

Age, gender, prior CLL duration, LDH value, BM histology, PB lymphocyte count, and the introduction of IFN-α in the induction regimen, showed no significant effect on the probability of achieving a response (CR+PR) while only 2 factors showed a significant and independent effect on the probability of achieving a response (CR+PR): the baseline initial Rai stage ($p = 0.05$) and the rate of lymphocyte reduction, ≤25% or >25%, recorded after the second course of therapy ($p = 0.01$) (Table 3).

**Post-induction phase**

Second randomized phase of the study: IFN-α vs no therapy. Seventy-eight of the 113 patients who had a response (69%) were subsequently randomized to receive maintenance treatment with IFN-α (arm C: 41 patients) or only clinical observation (arm C).

Table 3. Significant and independent prognostic factors for the probability of achieving a response, response duration and survival probability (in parentheses non-significant variables).

<table>
<thead>
<tr>
<th>Probability of achieving a response</th>
<th>Independent prognostic factors</th>
<th>p</th>
<th>CI     95%</th>
</tr>
</thead>
<tbody>
<tr>
<td>(Age, gender, Rai stage, prior CLL duration, LDH value, BM histology, PB lymphocyte count, induction therapy regimen, rate of peripheral blood lymphocyte reduction after the second course of induction therapy)</td>
<td>1st randomized phase (133 pts)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Probability of response duration</td>
<td>2nd randomized phase (78 pts)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Survival probability</td>
<td>1st randomized phase (133 pts)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(Gender, prior CLL duration, LDH value, BM histology, PB lymphocyte count, the induction therapy regimen, the quality of response, the rate of lymphocyte reduction, recorded after the second course of therapy)</td>
<td>2nd randomized phase (78 pts)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

In addition to the above mentioned variables, the administration of IFN-α as maintenance therapy
D: 37 patients). The baseline characteristics of patients were well balanced between the two treatment groups (Table 4). The reason for early (<6 months) IFN-α discontinuation in 8 patients (20%) was IFN-α-related toxicity including: neurotoxicity (5 patients), persistent febrile flu-like syndrome (2 patients) and persistent thrombocytopenia (1 patient). The reasons for late (>6 months) IFN-α discontinuation were a life-threatening car accident in 2 patients, a second malignancy (12%), acute myeloid leukemia (15%), infection (15%), Richter’s transformation (15%), second malignancy (12%), acute myeloid leukemia (5%) and myocardial infarction (2%). A patient included in arm A died during induction therapy because of pneumonia. The introduction of IFN-α at any time of the treatment approach was not significantly related to an increased mortality or to a specific cause of death. Nineteen patients who responded (17%) were excluded from the second randomization. The reasons for the protocol deviation in the second phase of the study were: one patient with residual marked splenomegaly underwent splenectomy, 14 young patients (median age 50 years) underwent an autologous stem cell transplantation and 4 patients refused the second randomization.

Survival and factors predicting survival
Causes of death. The median follow-up was 51 months. At the time of the analysis, 41 patients (31%) had died. The main cause of death was CLL progression (51%); other causes of death were infection (15%), Richter’s transformation (15%), second malignancy (12%), acute myeloid leukemia (5%) and myocardial infarction (2%). A patient included in arm A died during induction therapy because of pneumonia. The introduction of IFN-α at any time of the treatment approach was not significantly related to an increased mortality or to a specific cause of death.

Prognostic factors for survival
The overall actuarial survival probability at 6 years was 55%. The survival probability for the two induction arm groups was not significantly different (at 6 years, arm A: 57% vs arm B: 51%; p = 0.3). While gender, prior CLL duration, LDH value, BM histology, Rai stage, PB lymphocyte count, the rate of lymphocyte reduction, ≤25% or >25%, recorded after the second course of therapy and the type of the induction regimen, showed no significant effect on the response duration, a significantly longer response duration was shown by 2 groups of patients: patients randomized to receive IFN-α

### Table 4. Characteristics of patients included in the 2nd randomized phase of the study.

<table>
<thead>
<tr>
<th></th>
<th>No. of patients</th>
<th>IFN therapy</th>
<th>Clinical observation</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Median age</strong></td>
<td>55 (78%)</td>
<td>56</td>
<td>54 (37 pts. %)</td>
<td>NS</td>
</tr>
<tr>
<td><strong>Gender</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>58 (74)</td>
<td>33 (80)</td>
<td>25 (68)</td>
<td>NS</td>
</tr>
<tr>
<td>Female</td>
<td>20 (26)</td>
<td>8 (20)</td>
<td>12 (32)</td>
<td></td>
</tr>
<tr>
<td><strong>Time to therapy</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤1 yr</td>
<td>35 (48)</td>
<td>17 (41)</td>
<td>18 (49)</td>
<td>NS</td>
</tr>
<tr>
<td>&gt;1 yr</td>
<td>43 (52)</td>
<td>24 (59)</td>
<td>19 (51)</td>
<td></td>
</tr>
<tr>
<td><strong>Rai stage</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I-II</td>
<td>63 (81)</td>
<td>36 (88)</td>
<td>27 (73)</td>
<td>NS</td>
</tr>
<tr>
<td>III-IV</td>
<td>15 (19)</td>
<td>5 (12)</td>
<td>10 (27)</td>
<td></td>
</tr>
<tr>
<td><strong>Induction therapy</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>FLU + PDN</td>
<td>38 (49)</td>
<td>19 (46)</td>
<td>19 (51)</td>
<td>NS</td>
</tr>
<tr>
<td>FLU + PDN + IFN-α</td>
<td>40 (51)</td>
<td>22 (54)</td>
<td>18 (49)</td>
<td></td>
</tr>
<tr>
<td><strong>Response to induction</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CR</td>
<td>33 (42)</td>
<td>19 (44)</td>
<td>15 (41)</td>
<td>NS</td>
</tr>
<tr>
<td>PR</td>
<td>45 (58)</td>
<td>23 (56)</td>
<td>22 (59)</td>
<td></td>
</tr>
</tbody>
</table>

NS: not significant.
the second randomized phase of the study was separately analyzed, gender, prior CLL duration, LDH value, BM histology, PB lymphocyte count, the introduction of IFN-α in the induction regimen, the quality of response, the rate of lymphocyte reduction (≤25% or >25% recorded after the second course of therapy) and the administration of IFN-α as maintenance therapy showed no significant effect on survival duration, while, again, age (p = 0.01) and Rai stage (p = 0.01), emerged as significant and independent factors (Table 3).
Toxicity

The main toxicities observed during induction therapy were myelosuppression and infections which occurred with no significant differences in the two arms. After the 729 administered courses of fludarabine + prednisone ± IFN-α, we observed 46 cases (6%) of severe granulocytopenia (WHO grade III-IV), 7 episodes (0.9%) of fever of unknown origin (FUO) and 10 cases (1.4%) of pneumonia. A case of interstitial pneumonia with negative broncho-alveolar lavage, 2 cases of Listeria monocytogenes sepsis, 6 cases of dermatomal HVZ and 7 of herpes simplex were observed during induction therapy. In 2 patients, a reactivation of viral hepatitis (HBV) was observed. One patient had a cerebral hemorrhage after completion of therapy. During the first course of induction therapy 4 patients who received IFN-α in addition to fludarabine and prednisone reported flu-like symptoms.

Persisting cytopenia (granulocytopenia in 5 cases and anemia in 1 patient) and the occurrence of an IgM autoimmune hemolytic anemia in 1 patient after 5 courses of fludarabine and prednisone, were the reasons for exclusion from the second randomization for 7 patients. Toxicity in patients randomized to receive IFN-α as maintenance therapy has already been reported in the post-induction phase section. Among patients randomized to clinical observation, 1 developed an IgG autoimmune hemolytic anemia 10 months after discontinuation of fludarabine and prednisone and 2 developed dermatomal HVZ.

Richter’s syndrome, acute leukemia and second malignancies

No statistically significant differences in the rates of Richter’s syndromes and solid tumors were observed in patients who at any time of their treatment course received or not IFN-α. At a median time of 16 months (range: 10-26 months) after the start of therapy, a histologic diagnosis of Richter’s syndrome had been made in 8 patients (6%) and included 7 cases of non-Hodgkin’s lymphoma (nodal: 2 patients; gastric: 1 patient; BM: 1 patient; skin: 1 patient; otorhinolaryngologic: 1 patient; orbit: 1 patient) and 1 case of Hodgkin’s lymphoma (cervical nodes and mediastinum). An acute myeloid leukemia (AML) was diagnosed in 2 patients (1.5%) at 12 and 28 months after the start of therapy. AML was not considered directly related to the treatment study since in both cases it occurred after the start of a subsequent chemotherapeutic approach that included alkylating agents. The first patient, a 57-year-old female, was treated with fludarabine + prednisone and then with IFN-α which was discontinued after 5 months because of CLL progression. Six months after starting a second-line treatment with chlorambucil a diagnosis of AML was made. The second case of AML was observed in a 55-year old patient who after fludarabine and prednisone therapy, underwent peripheral blood stem cell mobilization with high dose cyclophosphamide and then an autologous stem cell transplantation which was followed 16 months later by AML associated with lung cancer.

Four patients developed a second cancer during the induction therapy (fludarabine plus prednisone: 2 patients; fludarabine plus prednisone and IFN-α: 2 patients). In 2 cases there was no evidence of the tumor (liver, lung) in the baseline radiographic exams, while in the other 2 cases, the tumor involved organs not explored during the pre-therapy work-up (stomach, larynx).

Five patients developed a second malignancy (kidney: 2 patients; bladder: 1 patient; colon: 1 patient; melanoma: 1 patient) at a median time of 26 months (range: 12-37 months) after the start of therapy. The concomitant occurrence of two malignancies during the follow-up was observed in 3 patients (lung carcinoma + AML; kidney carcinoma + NHL; bladder carcinoma + parotid adenoma).

Discussion

This study was carried out in previously untreated CLL patients with advanced disease to evaluate the therapeutic benefit of the combination of IFN-α with a regimen including fludarabine and prednisone, and the role of IFN-α as maintenance therapy in prolonging the response duration.

Our results indicate that no therapeutic benefit is obtained by the addition of IFN-α to the fludarabine and prednisone regimen, since no difference was observed in terms of response rate between the two induction arm groups, being 86% for patients treated with fludarabine and prednisone and 84% for those treated with the same schedule combined with IFN-α. The high overall response rates compare favorably with those previously reported by other studies in which fludarabine was administered as a single agent, associated with prednisone or with cyclophosphamide.

While the introduction of IFN-α in the induction regimen had no impact, Rai stage and the rate of lymphocyte reduction (≤ 25% or > 25% after the second course of therapy) emerged as significant and independent predictive factors of the probability of achieving a response to therapy. These findings suggest that patients with advanced stages of CLL and a less than 25% reduction of PB lymphocytes after two courses of therapy represent a subset of poor responders who should be shifted to an intensified therapeutic approach, such as fludarabine associated with synergistic drugs and/or monoclonal antibodies.

Patients randomized to receive IFN-α as maintenance therapy and patients who achieved a CR showed a significantly longer response duration.
However, the multivariate analysis showed that the quality of the response was the only significant and independent factor influencing the duration of response. This finding, also reported by other studies,\textsuperscript{3,37} shows that a prolonged response in CLL results from a good quality response.

The mechanism of action of IFN-α in CLL is not yet fully understood. Different effects have been ascribed to IFN-α: an influence on cell survival through a modulation of bcl-2 protein expression\textsuperscript{40,41} or through the reduction of tumor necrosis factor,\textsuperscript{42-44} and changes in expression of adhesion molecules influencing the recirculation and homing of B cells.\textsuperscript{45,46} Morabito et al.\textsuperscript{47} observed that IFN-α enabled CLL cells to increase their intravitro resistance to fludarabine–induced cell death in some samples, while in others it produced a synergistic effect with fludarabine. It has been assumed by several authors\textsuperscript{48-49} that the anti-tumor activity of IFN-α could be ascribed to the activation of natural killer and T-cell mediated cytotoxicity. Thus, the significant depletion of T lymphocytes, usually recorded in CLL patients treated with fludarabine or fludarabine and prednisone,\textsuperscript{35,37} may have hampered the activation of the immune-mediated cytotoxicity by IFN-α and this could help to explain the absence of a therapeutic effect of IFN-α during the induction therapy. On the other hand, the evidence of a therapeutic effect of IFN-α given as post-remission therapy could be explained by the restoration of the T-lymphocyte count and by the lower leukemic burden.

A beneficial effect of IFN-α given as maintenance therapy has been reported in CLL patients who had achieved a prior response to chlorambucil\textsuperscript{23} or fludarabine.\textsuperscript{24} Furthermore, Ferrara et al.\textsuperscript{50} also reported an advantage of interferon given as maintenance therapy in a study including 45 CLL patients randomized to receive IFN-α or no treatment after achieving a response to the MiNa protocol (vincristine, cyclophosphamide, melphalan, peptichemo, and prednisone). No benefits were reported by O’Brien et al.\textsuperscript{51} in 31 CLL patients treated with IFN-α after first-line or second-line therapy with fludarabine (30 patients) or with chlorambucil + prednisone (1 patient). It is worth noting that in the latter study, only patients who, after 6 months of IFN-α therapy, showed a tumor response or a stable disease with at least an improvement in some immunologic parameters continued the IFN-α therapy beyond 6 months while in our study IFN-α was given as maintenance therapy until disease progression. The longer duration of IFN-α administration could have favorably influenced the activity of IFN-α and the response duration.

However, IFN-α was not an easily maintenance therapy for a considerable proportion of CLL patients, since 20% of patients discontinued IFN-α early because of therapy-related side effects.

No benefits in terms of reduced infection-related mortality, morbidity and spectrum of infections could be ascribed to IFN-α administration. In this study, which included only untreated patients, the addition of steroids to fludarabine did not result in an increase of clinically significant atypical infections as had been previously observed in studies in which mainly pre-treated patients were enrolled.\textsuperscript{35} The occurrence of AML in 2 patients who received an alkylating agent following the fludarabine + prednisone regimen (in one case with IFN-α, in the other without) ± IFN-α therapy, suggests, as previously reported,\textsuperscript{52} that this type of sequential treatment could represent a leukemogenesis risk.

In conclusion, the results of the present study indicate that fludarabine + prednisone produce a high response rate in previously untreated CLL patients with advanced disease, while no benefit is derived from adding IFN-α to this regimen. Despite the more prolonged response obtained by administering IFN-α as maintenance therapy, the quality of response after induction therapy was the only significant and independent factor predicting a prolonged response. Taken together, these findings justify more cytoreductive induction approaches and suggest that IFN-α therapy has only a limited role in the post-remission management of responsive CLL patients.

References


Pre-publication report

Contributions and Acknowledgments
FM designed the study. FRM with PZ, FZ, MG, VS, LM, RF were responsible for the care of patients and data collection. MLV was responsible for the statistical analyses. FRM with PZ, FZ, MB, ST and FM contributed to revising the manuscript. The authors are listed according to a criterion of decreasing individual contribution to the work, with the exception of the last author who had a major role as senior author in interpreting the data. FRM, PLZ, FZ are the authors taking primary responsibility for the paper. All tables and figures were created by FRM and MG.

We thank Dr Fortunato Morabito for his contribution and useful suggestions and Dr Antonella De Pascale for her help in managing this study.

Disclosures
Conflict of interest: none.
Redundant publications: no substantial overlapping with previous papers.

Funding
This work was supported by Roma AIL (Italian Association Against Leukemia-Section of Rome and Udine).

Manuscript processing
This manuscript was peer-reviewed by two external referees and by an Associate Editor. The final decision to accept this paper for publication was taken by the Editors. Manuscript received June 30, 2003; accepted October 15, 2003.