Resistance to All-\textit{Trans} Retinoic Acid (ATRA) Therapy in Relapsing Acute Promyelocytic Leukemia: Study of In Vitro ATRA Sensitivity and Cellular Retinoic Acid Binding Protein Levels in Leukemic Cells

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All-\textit{trans} retinoic acid (ATRA) induces leukemic cell differentiation and complete remission (CR) in a high proportion of patients with acute promyelocytic leukemia (AML3 subtype). However, relapses occur when ATRA is prescribed as maintenance therapy, and resistance to a second ATRA-induction therapy is frequently observed. An induced hypercatabolism of ATRA has been suggested as a possible mechanism leading to reduced ATRA sensitivity and resistance. CRABPII, an RA cytoplasmic binding protein linked to RA’s metabolism pathway, is induced by ATRA in different cell systems. To investigate whether specific features of the AML3 cells at relapse could explain the in vivo resistance observed, we studied the CRABP levels and in vitro sensitivity to ATRA of AML3 cells before and at relapse from ATRA. Relapse-AML3 cells (n = 12) showed reduced differentiation induction when compared with "virgin"-AML3 cells (n = 31; P < .05). Dose-response studies were performed in 2 cases at relapse and showed decreased sensitivity to low ATRA concentrations. CRABPII levels and in vitro differentiation characteristics of AML3 cells before and at relapse from ATRA therapy were studied concomittantly in 4 patients. High levels of CRABPII (median, 20 fmol/mg of protein) were detected in the cells of the 4 patients at relapse but were not detected before ATRA therapy. Three of these patients showed a decrease in differentiation induction of their leukemic cells, and a failure to achieve CR with a second induction therapy of ATRA 45 mg/m²/day was noted in all patients treated (n = 3). Results from this study provide evidence to support the hypothesis of induced-ATRA metabolism as one of the major mechanisms responsible for ATRA resistance. Monitoring CRABPII levels after ATRA withdrawal may help to determine when to administer ATRA in the maintenance or relapse therapy of AML3 patients.

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induced during ATRA therapy and result in overall insufficient nuclear-ATRA levels that lead to reduced RA-sensitivity in vivo.

MATERIALS AND METHODS

Patients. During the period September 1990 to September 1991, 12 AML3 patients with the specific translocation t(15;17) were studied at relapse after ATRA-induced CR. First induction ATRA therapy (45 mg/m²/d) was administered to these patients until CR was obtained (median, 3 months). After achievement of CR, the patients were given low-dose maintenance chemotherapy (6-mercaptopurine/methotrexate n=11; cytarabine/methyl GAG n=1), and 1 patient was kept on ATRA alone. Relapse occurred after a median withdrawal interval from ATRA of 6 months (0 to 26 months) (Table 1). Six patients were treated with a second ATRA induction therapy (45 mg/m²/d). Failure, as noted by an absence of in vivo cytologic signs of maturation in the blood and bone marrow (BM) and by a rapid increase of blast cells leading to the addition of conventional chemotherapy, was followed by early death in 5 cases after a median therapy of 15 days.

Cell samples and culture. BM samples were collected after informed consent in heparinized tubes before the first ATRA therapy and when relapse occurred. Leukemic cells were isolated after adherence of monocytes to the plastic surface. Leukemic cells were maintained in suspension culture in culture medium consisting of RPMI 1640 (GIBCO, Paisley, UK) supplemented with 15% fetal calf serum (FCS; Seromed, Berlin, Germany), L-glutamine (2 mmol/L), penicillin (100 IU/mL), and streptomycin (1%; GIBCO), incubated at 37°C in a humidified atmosphere of 5% CO₂ in air, and subcultured twice a week. Cell number and viability were estimated by trypan-blue exclusion dye test.

The enriched leukemic cell fraction (>90% blasts) was suspended in culture medium at a concentration of 10⁶ cells/mL in the presence or absence of ATRA (provided by Dr W. Bollag, Hoffmann-La Roche, Basel, Switzerland). Cell aliquots were taken at days 3 and 5 or later for assessment of viability and differentiation status. Differentiation was assessed on morphologic and functional criteria. Slides were stained with May-Grünwald-Giemsa stain (Laboratoire Central d’Hématologie, Hôpital Saint-Louis). The nitroblue tetrazolium (NBT) reduction test was performed using standard methodology with phorbol myristate acetate (PMA) as previously described.38 Differential counts were performed under light microscopy on a minimum of 200 cells. The percentage of cells containing intracellular-reduced black formazan deposits was thus determined.

Preparation of cytosolic extracts. Fresh human leukemic cells (30 x 10⁶) were washed once in medium without FCS and in cold sodium phosphate buffer before protein extraction. Cells were homogenized at 4°C in 500 µL of an ice-cold extraction buffer (Tris, 100 mmol/L; NaCl, 50 mmol/L; EDTA, 5 mmol/L; dithiothreitol, 2 mmol/L; pH 7.4) with three 30-second strokes of a Polytron (Kinematica; GiM, Luzern, Switzerland) at full speed. Supernatants were obtained by centrifugation at 100,000g at 4°C for 60 minutes, distributed in 100 µL aliquots, and frozen at -20°C or used immediately.

Incubation of cytosolic extracts with ATRA. All manipulations were performed under dim or yellow light. Supernatant aliquots of 100 µL containing 300 µg protein were incubated with 300 nmol/L of [³H]-ATRA (specific activity, 2,035 GBq/mmol) from DuPont New England Nuclear (Boston, MA) at 4°C for 16 hours. For binding specificity a 200-fold excess of unlabeled ATRA (Sigma Chemical Co, St Louis, MO) was added in the same manner as for the radioactive compounds.

Slab polycrylamide gel electrophoresis (PAGE). At the end of the incubation period, an aliquot of the supernatant (70 µL) was subjected to vertical slab PAGE under dim light at 14°C as described by Siegenthaler et al with slight modifications of the method.32,39 The proteins were separated according to their different electrophoretic mobilities in the gel, as a consequence of their net charge. This technique allows the separation of CRABPI and II. The bands were collected and treated overnight with 400 µL Protosol (DuPont, Boston, MA) in 5-mL scintillation vials before 4 mL of solubilization cocktail was added. The radioactivity was then determined in a scintillation counter.

Statistical analysis. Differences in the percentage of differentiated cells at onset and at relapse were assessed by the parametric Sign test.40

RESULTS

In vitro differentiation of AML3 cells from patients relapsing from ATRA therapy. In vitro response to ATRA 10⁻⁶ mol/L was studied in 12 AML3 patients who relapsed after ATRA therapy (45 mg/m²/d). Table 1 summarizes the clini-
cal characteristics of these patients before relapse occurred, namely length of ATRA therapy and of the disease-free survival under low-dose maintenance chemotherapy. All patients at relapse after ATRA therapy had a cytologic and/or cytogenetic diagnosis of AML3.

The purified leukemic cells were cultured for 5 days in the presence of ATRA $10^{-6}$ mol/L. Compared with 31 virgin AML3 cell samples studied during the same period, relapse cases achieved a lower percentage of differentiated cells (median 46% vs 90% NBT positive cells, $P < .05$; Fig 1A). Eight of the 12 relapse cases (case nos. 1 through 7 and 11) and only 4 of 31 virgin cases showed less than 70% differentiated cells using the oxidative burst function criteria (Fig 1A). Three of the relapse cases had less than 35% differentiated cells (case nos. 2, 3, and 5), and one case did not respond (case no. 1). Absence of morphologic modifications correlated with the poor functional achievement. Cell count was similar in these groups (med $1 \times 10^6$ cells/mL in virgin AML3 samples vs $0.8 \times 10^6$ cells/mL at relapse). Therefore, the failure of differentiation could not be due to an increased proliferation of the undifferentiated AML3 cells. Virgin AML3 cells are extremely sensitive to ATRA, with 80% of the cell population already reducing NBT $10^{-7}$ mol/L and 45% at $10^{-8}$ mol/L (Fig 1A and B). Sensitivity to low ATRA concentrations was studied in 2 AML3 samples at relapse after ATRA therapy (case nos. 2 and 10). In both cases, absence of sensitivity to ATRA concentrations of $10^{-7}$ mol/L and $10^{-8}$ mol/L was observed (Fig 1B).

The in vitro response to ATRA $10^{-6}$ mol/L of 12 AML3 patients at relapse was compared with the differentiation achieved on the AML3 cells of these same 12 patients before any ATRA therapy. A decrease in the percentage of differentiated cells in the leukemic cell culture was clear in half of the cases ($P < .03$; 5 cases [nos. 1 through 5] with a reduction of at least 23% at $10^{-6}$ mol/L of ATRA; Fig 2A). Low dose-response analysis was possible in 1 patient (case no. 10) before ATRA therapy and when relapse occurred. A shift to the right of the dose-response curve at relapse was observed (Fig 2B). These data suggest that a decrease in ATRA sensitivity may not be detected in all AML3 cell samples at relapse with a saturating $1 \mu$mol/L concentration of ATRA and that lower doses should be tested at onset and at relapse.

In 4 cases (nos. 2, 4, 5, and 10), higher ATRA concentrations were studied to test the hypothesis that relapse-AML3 cells could be triggered to differentiate with higher ATRA concentrations. Effective restoration of differentiation was only observed in 1 case (no. 2) after incubation with $10^{-5}$ mol/L ATRA. Cases no. 4 and 10 already had a relatively good response at $10^{-6}$ mol/L and case no. 5, which had a poor response, was not improved by higher RA concentrations (Table 2).

**Presence of CRABPII in relapse-AML3 cells.** In a previous study, we reported the induction of the CRABPII in normal hematopoietic cells of AML3 patients undergoing ATRA therapy (Figs 3A and B; and Figs 4A, B-1, and B-2). CRABPII is distinguished from CRABPI by a different binding affinity for retinoids and a different isoelectric point detected by PAGE. Cytosol extracts of 4 AML3-
Fig 2. (A) In vitro differentiation of AML3 cells of 12 patients with ATRA 10^-6 mol/L before and after ATRA therapy at relapse (expressed as the percentage of NBT positive cells at relapse minus the percentage at onset of ATRA therapy). (B) Dose-effect of ATRA on patient no. 10 AML3 cells at onset (C) and at relapse after ATRA therapy (W).

cell samples at onset and at relapse (patients no. 2, 4, 5, and 10) were incubated with [3H]-ATRA and subjected to vertical PAGE. The purified leukemic cell samples from these patients averaged 74.5% and 67.4% blasts at onset and at relapse, respectively (Table 1). A specific radioactive peak of [3H]-ATRA was detected between fraction 18 and 22 of migration and abolished by incubation with 200-fold excess of unlabeled ATRA (Fig 3D, patient no. 2). This peak eluted at the fractions corresponding to the CRABPII. The amount of CRABPII levels detected in the 4 relapse-AML3 cells (Fig 4C) were above the amounts observed in normal BM cells of ATRA-treated patients (Fig 3B, Fig 4B-1 and B-2). The exact function of this protein is not known, but recent data point to its role in the transport of ATRA to its metabolizing enzymes, thus acting as a modulator of intracellular ATRA concentrations.26 In F9 teratocarcinoma cells, reduced sensitivity to ATRA is correlated with decreased ATRA intracellular concentrations and with high levels of CRABP.28 The effect of ATRA in AML3 cells is exquisitely dose-dependent whether for induction of cell differentiation in vitro,38 for PML/RARα transactivation of ATRA-inducible genes,4 or for normal RARα induction in AML3 cells.31 Thus, it may be expected that variations in ATRA concentrations may result in reduced biological response. The presence of high levels of CRABPII in relapse-AML3 cells could reduce the effective cellular concentration of ATRA and may result in reduced differentiation. However, in the present study, a decrease in the sensitivity of AML3 cells in vitro to 10^-6 mol/L ATRA was observed in only half of the cases. In vitro response could have predicted in vivo failure in only 4 of 6 cases, whereas in virgin AML3 cells, in vitro response has always been correlated with in vivo remission.38,42

Table 2. Dose-Effect of ATRA Differentiation in Relapse-AML3 Cells

<table>
<thead>
<tr>
<th>Patient No.</th>
<th>Onset</th>
<th>Relapse</th>
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<tr>
<td></td>
<td>Control (mol/L)</td>
<td>Control (mol/L)</td>
</tr>
<tr>
<td>2</td>
<td>3</td>
<td>82</td>
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<tr>
<td>4</td>
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<td>58</td>
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<tr>
<td>10</td>
<td>12</td>
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Values are the percentage of differentiated cells after 5 days of in vitro culture with ATRA.

Abbreviation: ND, not determined.
Two explanations may be forwarded for this apparent discrepancy between in vitro and in vivo response at relapse. The first is that the concentration of ATRA prevailing in most in vitro differentiation assays (10^{-7} mol/L) is above the maximal effective concentration in the AML3 system. We have previously stressed the potency of a 10^{-7} mol/L concentration to induce maximal differentiation in AML3 cells. A decrease in ATRA-induced differentiation might have been observed in all relapse cases if nonsaturating concentrations (10^{-7} mol/L) had been used. In fact, in the only 2 cases in which this could be tested, we were able to show a clear shift to the right of the dose-response curve at low ATRA concentrations, suggesting that AML3 cells at relapse may well have a reduced sensitivity to ATRA. To study this hypothesis, in vitro models are being set up in the laboratory.

The second explanation is that poor in vivo response may not necessarily be attributed to the characteristics of the relapsing leukemic cell alone. Low ATRA plasma levels have been reported during ATRA therapy. We have previously linked low plasma ATRA concentrations to increased clearance rate and poor in vivo response. An overall increase of ATRA catabolism in cell tissues by an increase of CRABP, as it has been shown in the skin and hematopoietic tissue, may account for the low plasma ATRA concentrations observed during continuous ATRA therapy and for failure to achieve effective ATRA intracellular concentrations.

Thus, one may hypothesize that, in AML3 patients, ATRA therapy induces a cascade of events. The aim of one of these is to subsequently reduce the intracellular concentration of ATRA. Increase in the level of proteins linked to ATRA's metabolism during ATRA therapy is one major step in this process. Induction of CRABPII in AML3 patients may result from a direct regulation by ATRA, because an RA-responsive element has been reported in the promoter region of the mCRABPII gene. Along with the increase of CRABP, increased levels of cytochrome P450 enzymes have been observed in ATRA-treated hamsters and linked to the hypercatabolic state observed in AML3 patients treated with ATRA. For some of these patients, we have different data showing that, at one point during CR, CRABP levels were no longer detectable. This suggests that a residual leukemic clone that was CRABP positive during ATRA therapy persisted at CR. Furthermore, the fact that high levels of CRABP are detected both in the normal and leukemic BM cells of AML3 patients during ATRA therapy suggests that the hypercatabolic state may be responsible both for the relapses observed if ATRA is maintained as consolidation therapy and for the subsequent resistance to ATRA when relapse occurs.

The majority of the patients in this early clinical trial (1990) were either still under ATRA therapy or after a short withdrawal period from ATRA when they relapsed, forwarding an explanation for the presence of CRABPII in the leukemic cells. CRABPII levels decrease slowly (median, 1 month) after ATRA withdrawal. Therefore, it may be expected that the hypercatabolic state may eventually be reduced after ATRA withdrawal and sensitivity to ATRA restored. In fact, 2 relapse-AML3 patients have been recently shown to respond to a second ATRA therapy, after withdrawal period of 6 months and 2 years (S. Castaigne, and R.P. Warrell Jr, personal communications, 1992). Although one of the patients in this study relapsed and failed to respond to ATRA after a withdrawal period of 10 months, one must keep in mind that most of the patients in this study had already relapsed after previous chemotherapy before receiving their first induction ATRA therapy and may have had other factors leading to reduced RA sensitivity in vitro and in vivo. It is equally important to stress that, although the increased ATRA metabolism may be acquired in all AML3 patients undergoing ATRA therapy, alternative or coexisting factors of resistance, such as molecular changes and selection of non-AML3 clones, cannot be ex-
Fig 4. CRABP II protein level in normal and leukemic hematopoietic cells of AML3 patients undergoing ATRA therapy. AML3 cells before therapy (A); during ATRA therapy at 60 days (B-1); at 90 days (B-2), and after 1 month of withdrawal (B-3); and in relapse (C) after ATRA therapy (withdrawal time of a maximum of 10 months).

clined and have occasionally been observed (C.C., personal observations, 1993).

Recent results from our study group suggest that maintenance therapy in AML3 could be improved if ATRA was combined with chemotherapy. Thus, it is necessary to overcome the inconvenience of induced RA resistance. Strategies to decrease the induction of the hypercatabolic state (lower ATRA concentrations, shorter induction regimens, choice of new retinoids, or metabolizing, limiting drugs such as ketonozole) are desirable. Monitoring CRABP levels by PAGE or reverse transcriptase-polymerase chain reaction (undergoing study) after ATRA withdrawal may determine the best time and schedule to readminister ATRA in the maintenance or relapse therapy of AML3 patients.

REFERENCES


Resistance to all-trans retinoic acid (ATRA) therapy in relapsing acute promyelocytic leukemia: study of in vitro ATRA sensitivity and cellular retinoic acid binding protein levels in leukemic cells [see comments]

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