

ACUTE PROMYELOCYTIC LEUKEMIA COMPLICATING CHEMO-RADIOTHERAPY FOR MULTIPLE MYELOMA

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A 61-year-old woman was admitted to our Department for severe gastrointestinal bleeding. She had been diagnosed with IgA κ multiple myeloma three years earlier and had been treated with both chemotherapy and radiotherapy (this latter for spinal cord compression by extramedullary myelomatous tissue).

On admission, gastroscopy showed a bleeding gastric ulcer and blood cell counts were the following: Hb 4.2 g/dL, WBC $1.9 \times 10^9/L$ (differen-

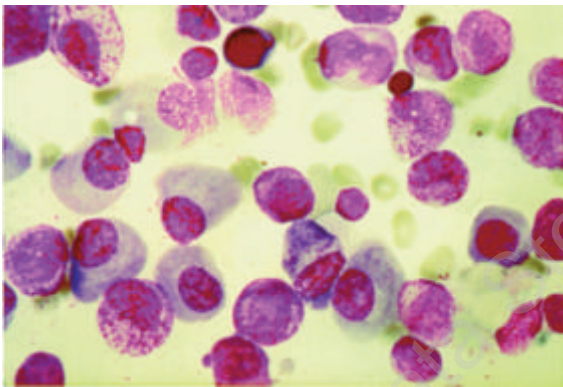


Figure 1. Bone marrow aspirate showing many plasma cells and abnormal promyelocytes. MGG x 640.

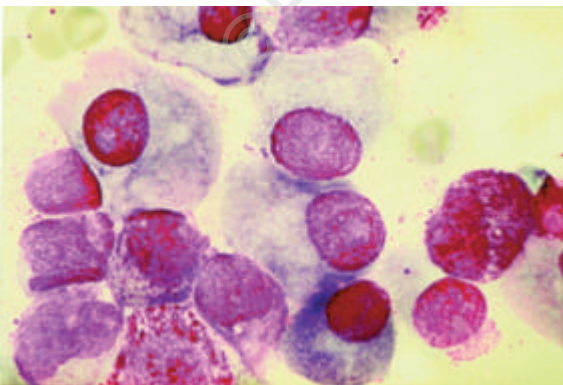


Figure 2. Higher magnification shows a promyelocyte and a cluster of plasma cells. MGG x 1200.

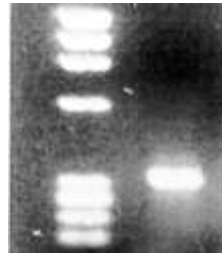


Figure 3. Detection of PML-RAR α fusion by RT-PCR in a bone marrow sample (right lane; left lane shows molecular weights expressed in base pairs). The ethidium bromide-stained gel shows a 326-bp band corresponding to the BCR1 PML-RAR α junction.

tial count: neutrophils 43%, eosinophils 4%, basophils 1%, lymphocytes 47%, monocytes 5%), and platelets $9 \times 10^9/L$. Bone marrow was heavily infiltrated by abnormal plasma cells, and abnormal promyelocytes were also found (Figures 1 and 2). Within 48 hours of admission PML-RAR α gene rearrangement¹ was identified in the patient's bone marrow by RT-PCR (Figure 3) and a diagnosis of secondary acute promyelocytic leukemia (APL) was formulated.

Treatment with all-trans retinoic acid (ATRA) induced a partial, transient remission; however, the woman eventually died of sepsis. APL has seldom been observed as a secondary malignancy;² most of the acute leukemias complicating antineoplastic chemotherapy and/or radiotherapy are usually either myelomonocytic or unclassifiable. Albeit for only a short period of time, the patient clearly benefitted from treatment with ATRA, which was essentially delivered on the basis of the molecular diagnosis.

References

1. Diverio D, Riccioni R, Mandelli F, Lo Coco F. The PML/RAR α fusion gene in the diagnosis and monitoring of acute promyelocytic leukemia. *Haematologica* 1995; 80:155-60.
2. Amadori S, Papa G, Anselmo AP, Fidani P, Mandelli F, Biagini C. Acute promyelocytic leukemia following ABVD (doxorubicin, bleomycin, vinblastine, and dacarbazine) and radiotherapy for Hodgkin's disease. *Cancer Treat Rep* 1983; 67:603-4.