

Figure 1. Light scatter properties of analyzed cells (top). The flow cytometric dot plots clearly show that virtually all CD19+ cells are positive for CD5 antigen and there are two cell populations with different HLA-DR antigen expression pattern. CD33 antigen is found to be the only antigen that expressed more than 50% of the cells and most of them are negative for HLA-DR antigen.

nosis but we do not have any doubts about the diagnosis because more then  $10\times10^{9}/L$  cells expressed CD5, CD19, CD20 and CD22 (Figure 1).

The concomitant presentation of AML and CLL is extremely rare and the use of two-color flow cytometry to differentiate the cell populations demonstrates the utility of this technology in the diagnosis of unusual hematologic malignancies.

Mustafa Nuri Yenerel,\* Ibrahim Hatemi,° Hüseyin Keskin\*

\* Istanbul University, Istanbul Medical School, Department of Internal Medicine, Division of Hematology, Çapa, Istanbul; " Haseki State Hospital, Haseki, Istanbul, Turkey

## Key words

CCL, AML, flow cytometry.

### Correspondence

Mustafa Nuri Yenerel, MD, Istanbul University, Istanbul Medical School, Department of Internal Medicine, Division of Hematology, Çapa, Istanbul, Turkey. Fax international +90.212.6311263.

#### References

 Caballero MD, Gonzalez M, Canizo MC, Orfao A, Nieto MJ, San-Miguel JF. Concomitant chronic lymphocytic leukemia (CLL) and acute myeloid leukemia. Complete remission of CLL achieved with high-dose cytosine arabinoside. Leukemia 1992; 6:856-8.

 Conlan MG, Mosher DF. Concomitant chronic lymphocytic leukemia, acute myeloid leukemia, and thrombosis with protein C deficiency. Case report and review of the literature. Cancer 1989; 63:1398-401.

 Rai KR, Patel DV. Chronic lymphocytic leukemia. In Hoffman R, Benz EJ J, Shattil SJ Furie B, Cohen HJ, Silberstein LE(eds): Hematology. Basic Principles and Clinical Practice. 2nd ed. Curchill Livingstone, New York, 1995, p 1308.

 Lima M, Porto B, Rodrigues M, et al. Cytogenetic findings in a patient presenting simultaneously with chronic lymphocytic leukemia and acute myeloid leukemia. Cancer-Genet Cytogenet 1996; 87:38-40.

 Mateu R, Bellido M, Sureda A, et al. Concomitant chronic lymphocytic leukemia and acute myeloid leukemia with an uncommon immunophenotype. Am JHematol 1997; 56:281.

 Tamul KR, Meyers DC, Bentley SA, Folds D. Two color flow cytometric analysis of concomitant acute myeloid leukemia and chronic lymphocytic leukemia. Cytometry 1994; 18:30-4.

# Acute megaloblastic anemia: homocysteine levels are useful for diagnosis and follow-up

Sir.

Vitamin B<sub>12</sub> (cobalamin) and folic acid deficiencies lead to megaloblastic anemia (MA), and induce accumulation of methylmalonic acid (MMA) and homocysteine (HCY). The most common presentation of MA is classical macrocytic anemia. Other presentations are acute megaloblastosis (AM) and masked megaloblastosis. This report, we present a case of AM diagnosed and followed up by evaluation of HCYlevels.

A 45-year old male was diagnosed as having Philadelphia-positive chronic myelogenous leukemia. Three years after diagnosis the patient developed a lymphoid blast crisis and was started on a chemotherapy protocol. The first consolidation treatment consisted of 6-mercaptopurine, methotrexate (MTX), VM-26 and cytarabine. MTX rescue with folinic acid was performed following standard guidelines. On day +14 a platelet count of 9×109/L was found. Hb was 99 g/L, mean corpuscular volume (MCV) 92 fL and leukocyte count was 7.06×109/L with 84% of neutrophils with hypersegmentation. Reticulocyte count was  $0.053 \times 10^{12}/L$  (1.66%). Vitamin B<sub>12</sub> levels and red cell folate were 322 pmol /L (normal 150-1200) and 938 nmol/L (normal 441-1285), respectively. A BM aspirate revealed 30% of erythroid precursors with megaloblastic features and a 55% of myeloid precursors with increased size and no blast cells. Serum HCYlevelswere 38 µmol/L (normal < 16). The

Haematologica vol. 84(8): August 1999



Table 1. Evolution of analytical parameters during folinic acid and vitamin B<sub>12</sub> treatment.

	Pre-treatment Day -9	Onset Day 0	Post-treatment Day +9
Platelets (x10º/ L)	134	9	112
Leukocytes (x109/L)	6.76	7.06	5.72
Hemoglobin (g/ L)	91	99	95
MCV(fL)	93	92	95.3
Reticulocytes (x1012/ L)	0.037	0.053	0.163
Homocysteine (µmol/L)	-	38	9

AM, acute megaloblastosis; MCV, mean corpuscular volume.

patient was diagnosed as having AM and began treatment with folinic acid 12 mg ivin one single dose and folic acid 5 mg/day po for 14 days and parenteral vitamin B<sub>12</sub> 2 mg/day for 4 consecutive days. After 10 days of treatment the platelet count increased to 112×109/L and reticulocyte count to 0.163×1012/L (5.41%). Vitamin B<sub>12</sub> level was 716 pmol/L, red cell folate level 1,506 nmol/L and serum HCY level decreased to normal value (9 µmol/L)

Four different clinical forms of megaloblastosis have been described.3,4 The classical form has an insidious onset with frequent neurologic symptoms and macrocytic anemia. Vitamin B<sub>12</sub> and/or red cell folate levels are decreased. The second form is the subtle MA anemia with ill-defined clinical symptoms and decreased or borderline vitamin B<sub>12</sub> and folic acid levels with other abnormalities (dUST, HCY, MMA).2 Masked megaloblastosis coexists with other deficiencies; MCV is normal or decreased. 56 MA of acute onset is the rarest form.3 There are two clinical presentations; the masked undiagnosed classical MA with cytopenias of abrupt onset and the so-called AM.3-7 In AM severe thrombocytopenia develops in 1 to 3 weeks, MCV is normal or only moderately increased. This presentation is more frequent in patients with risk factors: parenteral nutrition, infection, dialysis or treatment with some antifolate drugs. Mortality is high.3 The reticulocyte count is low. Vitamin B<sub>12</sub> and red cell folate levels are normal. BM aspirate shows megaloblastic changes. Classically, dUST is used as a diagnostic test. Nevertheless, HCY serum assays provide a sensitive test for the diagnosis of AM, especially in its early stages.8 In vitamin B<sub>12</sub> deficiences both HCY and MMA levels are high. In

folate deficiencies only HCY concentration is increased. 9,10 HCYlevels are also useful for AM followup of AM; levels return to normal after starting treatment with vitamin B<sub>12</sub> or folic acid. The evaluation of serum HCYlevels is an easy and non-invasive test for the diagnosis and follow-up of AM.

> Marina Carrasco, Angel Remacha, Anna Sureda, Filar Sardà, Rodrigo Martino, Jorge Sierra.

Department of Hematology, Hospital de la Santa Creu i Sant Pau, Barcelona, Spain.

# Key words

Acute megaloblastosis, folicadid, cobalamin, homooysteine

### Correspondence

Angel Remacha Sevilla, MD, Laboratorio de Hematología, Hospital de la Santa Creu i Sant Pau, Antoni Maria i Claret, 167, 08025 Barcelona, Spain. Phone: international +34-93-2919290 - Fax international +34-93-2919192 - Email: 2107@hsp.santpau.es

### References

- Green R. Metabolite assay in cobalamin and folate
- deficiency. Bailliére Clin Haematol 1995; 8:533-66. Carmel R. Subtle cobalamin deficiency. Ann Intern Med 1996; 124:338-40.
- Remacha A, Gimferrer E. Las megaloblastosis agudas: revisión y reconsideración conceptual de las distintas formas de presentación de las megaloblastosis. Biol Clin Hematol 1984; 6:167-82
- Carmel R. Pernicious anemia. The expected findings of very low serum cobalamin levels, anemia, and macrocytosis are often lacking. Arch Intern Med 1988; 148:1712-4
- Spivak J. Masked megaloblastic anemia. Arch Intern
- Med 1982; 142:2111-4. Bennett M, Koren A, Ludacer E. B12 deficiency in αthalassemia. N Engl JMed 1984; 310:1058-9.
- Martinez E, Remacha A, Roca-Cusachs A. Acute exacerbation of folate-dependent chronic megaloblastosis. Biol Clin Hematol. 1992; 14:223-9.
- Vester B, Rasmussen K. High performance liquid chromatography method for rapid and accurate determination of homocysteine in plasma and serum. Eur J
- Clin Chem Clin Biocherm. 1991; 29:549-54. Allen RH, Stabler SP, Savage DG, Lindenbaum J Diagnosis of cobalamin deficiency: I: usefulness of serum methylmalonic acid and total homocysteine concentrations. Am JHematol 1990; 34: 90-8
- 10. Lindenbaum J Savage DG, Stabler SP, Allen RH. Diagnosis of cobalamin deficiency. II: relative sensitivities of serum cobalamin, methylmalonic acid and total homocysteine concentrations. Am J Hematol 1990; 34:99-107.

Haematologica vol. 84(8): August 1999

