- Stasi R, Stipa E, Masi M, et al. Antiphospholipid antibodies: prevalence, clinical significance and correlation to cytokine levels in acute myeloid leukemia and non Hodgkin's lymphoma. Thromb Haemost 1993; 70: 568-72.
- Asherson RA, Block S, Houssiau FA, Hughes GRV. Systemic lupus erythematosus and lymphoma: association with an antiphospholipid syndrome. J Reumatol 1991; 18:277-9.
- Keung YK, Cobos E, Meyerrose GE, Roberson GH. Progressive thrombosis after treatment of diffuse large cell non Hodgkin's lymphoma and concomitant lupus anticoagulant. Leuk Lymphoma 1996; 20:341-5.
 Esmon NL, Smirnov MD, Esmon CT. Lupus anticoag-
- 8. Esmon NL, Smirnov MD, Esmon CT. Lupus anticoagulants and thrombosis: the role of phospholipids. Haematologica 1997; 82:474-7.

New technology and changing parameters of leukapheresis for blood cell transplantation

Sir

Clinical investigators have recently developed an innovative technique of leukapheresis (LK) for blood cell transplantation (BCT) referred to as AutoPBSC System. This technique offers the following advantages: a) better collection efficiency of CD34+ hematopoietic progenitor cells; b) reduced collection of platelets; c) higher quality of LK components in terms of reduced contamination by granulocytes, platelets, and erythrocytes; d) reduced LK volume; and e) automation.1 These advantages prompted us to evaluate the effectiveness of the AutoPBSC System and to extend classic parameters for starting LK, i.e., CD34+ cells $\geq 20/\mu L$ and platelets $\geq 30 \times 10^3/\mu L$, also to poor-mobilizer and/or thrombocytopenic patients, i. e., with CD34+ cells $\leq 20/\mu L$ and/or platelets $\leq 30 \times 10^3/\mu L$, respectively. We confirm the advantages of the AutoPBSC System and demonstrate that efficient LK can successfully be performed also in these categories of patients.

Ninety-six leukaphereses were carried out in 65 consecutive patients undergoing BCT for treatment of poor prognosis malignancies (13 multiple myeloma, 12 breast cancer, 8 Ewing's sarcoma family of tumors, 9 non-Hodgkin's lymphoma, 7 Hodgkin's disease, 6 ovarian cancer, 3 rhabdomyosarcoma, 3 desmoplastic small cell tumor, 1 Wilms' tumor, 2 non-small cell lung cancer, 1 yolk sac tumor). The LK procedure implied processing 2.5-fold the individual's blood volume and adaptation of the AutoPBSC software default and harvest frequency as described by Ravagnani *et al.*¹ At the time of LK, the mean CD34⁺ cell count per μ L was 106, the median 53, and the range 3-626; mean platelet count was $104 \times 10^3 \mu$ L, median 92×10^3 , range $15-456 \times 10^3$.

As detailed in Figure 1, the collection target of CD34+ cells $\geq 5 \times 10^6$ in a single LK was achieved in 100% (48/48) of procedures when the initial CD34+ cells $\geq 50/\mu$ L and 21% (10/48) when CD34+ cells $\leq 50/\mu$ L. Single LK in poor-mobilizer patients (n = 17) with CD34+ cell counts >10/ μ L and $\leq 20/\mu$ L (n=14) and $\leq 10/\mu$ L (n=15) yielded mean 2.1×106, median 1.8, range 0.7-3.7×106 and mean 1.7×106 CD34+ cells/kg, median 1.7, range 0.8-3.4×106, respectively; in

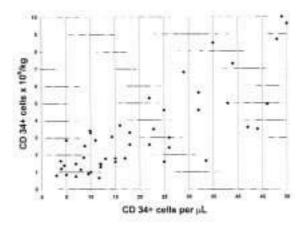


Figure 1. Yields of CD34⁺ progenitor cells by single LK versus CD34⁺ cell counts. Data shown are limited to the 48 procedures with CD34⁺ cells ≤50/µL (total LK=96).

thrombocytopenic patients (n = 7) a single LK yielded mean 3.9×10^6 CD34+ cells/kg, median 2.6, range $0.8\text{-}14.6\times10^6$; and in thrombocytopenic and poor-mobilizer patients (n = 5) it yielded mean 1.9×10^6 CD34+ cells/kg, median 1.8, range $0.8\text{-}3.1\times10^6$. Although platelet depletion in thrombocytopenic patients was negligible, a prophylactic platelet transfusion was given after LK to 3 patients.

Results of a single leukapheresis presented here compare favorably with those previously attained with other techniques^{1,2} and confirm for the first time the advantages of the AutoPBSC System in poor-mobilizer and thrombocytopenic patients as well, thus facilitating the clinical application of blood cell transplantation.

Katharina Granzow, * Roberta Schiavo, ° Inna Timofeeva, ° Gianalessandro Moroni, * Armando Santoro, ° Salvatore Siena °

*Servizio di Immunoematologia e Medicina Trasfusionale, Ospedale San Paolo, Milan; "Department of Oncology and Hematology and Section of Blood and Marrow Transplantation, Istituto Clinico Humanitas, Rozzano (MI), Italy

Key words

Leukapheresis, blood cell transplantation, CD34+ cells

Correspondence

Salvatore Siena, Divisione Oncologia Medica Falck, Ospedale Niguarda-Ca' Granda, piazza Ospedale Maggiore 3, 20162 Milan, Italy. Phone international +39.02.64442991 — Fax. international +39.02.64442910 — E-mail: salsiena@tin.it

References

- Ravagnani F, Siena S, De Reys S, et al. Improved collection of mobilized CD34+ hematopoietic progenitor cells by a novel automated leukapheresis system. Transfusion 1999; 39:48-55.
- Moog R, Muller N. Technical aspects and performance in collecting peripheral blood progenitor cells. Ann Hematol 1998; 77:143-7.