



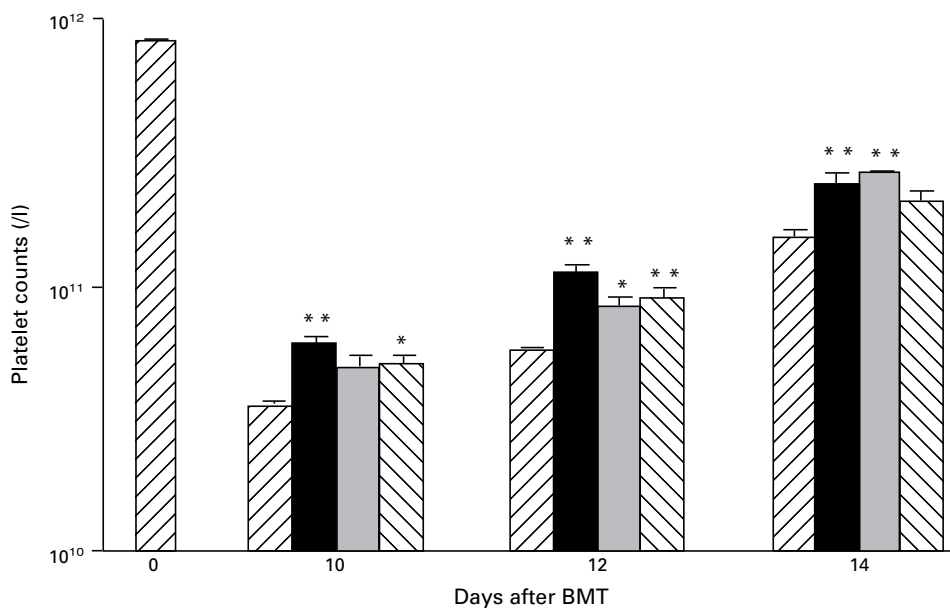
## Erratum

# Improvement of thrombocytopenia following bone marrow transplantation by pegylated recombinant human megakaryocyte growth and development factor in mice

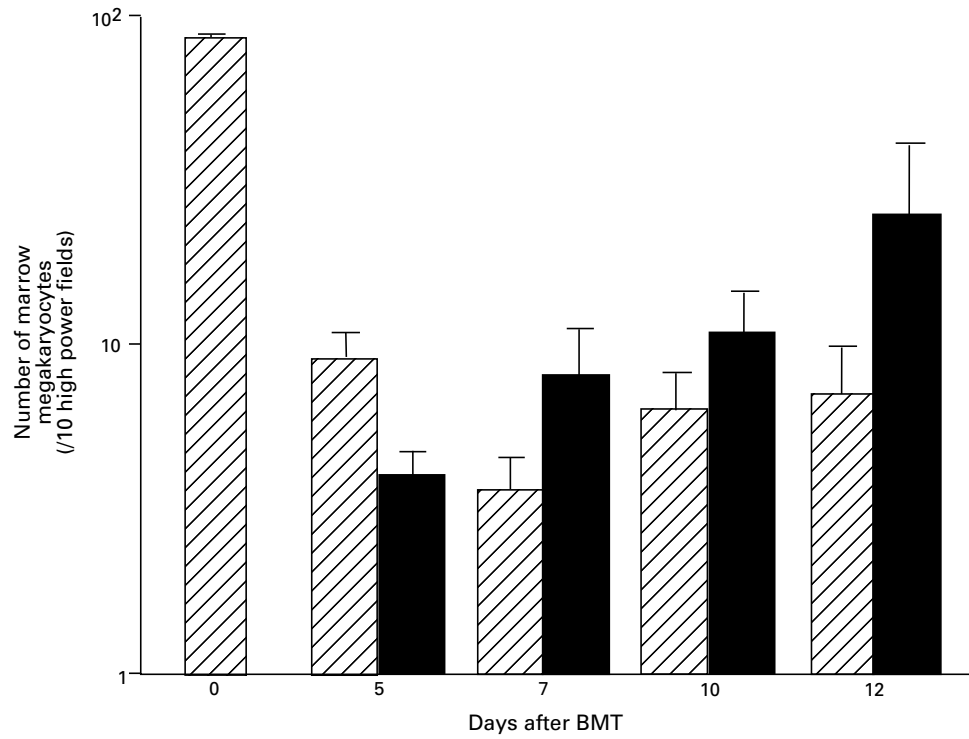
K Kabaya, K Shibuya, Y Torii, Y Nitta, M Ida, H Akahori, T Kato, M Kusaka and H Miyazaki

*Bone Marrow Transplantation* 1996; **18**: 1035-1041

In Figures 3 and 4 of the above paper some additional asterisks appeared in the final published copy of the paper. Corrected versions of the figures follow.



**Figure 3** Administration schedules of PEG-rHuMGDF following BMT in mice. Subcutaneous administration schedules of PEG-rHuMGDF were shown as follows: (1) consecutive treatment at a dose of 30  $\mu\text{g}/\text{kg}/\text{day}$  from the day after BMT (day 1) for 13 consecutive days (■); (2) alternate-day treatment at a dose of 55.7  $\mu\text{g}/\text{kg}/\text{day}$  on days 1, 3, 5, 7, 9, 11 and 13 (▨); and (3) at an interval of 3 days at a dose of 78  $\mu\text{g}/\text{kg}/\text{day}$  on days 1, 5, 9 and 13 (▧). The same volume of vehicle solution was injected as control from the day after BMT for 13 consecutive days (▩). Each bar indicates the mean  $\pm$  s.e. of five to six mice. Significant differences (\* $P < 0.05$ , \*\* $P < 0.01$ ) compared with the vehicle-treated mice were tested by Dunnett's test.



**Figure 4** Effects of PEG-rHuMGDF on the number of marrow megakaryocytes following BMT in mice. PEG-rHuMGDF at a dose of 30  $\mu\text{g}/\text{kg}/\text{day}$  (■) or vehicle solution (▨) was administered subcutaneously from the day after BMT for 13 consecutive days. Each bar indicates the mean  $\pm$  s.e. of three to five mice.