

## **ERRATUM**

In the article by Basser et al entitled "Randomized, Blinded, Placebo-Controlled Phase I Trial of Pegylated Recombinant Human Megakaryocyte Growth and Development Factor With Filgrastim After Dose-Intensive Chemotherapy in Patients With Advanced Cancer," which appeared in the May 1, 1996 issue (Vol 89, No 9, pp 3118-3128), corrections should be noted. In Table 5 the numbers were misaligned and figure legends 1 and 2 contained errors. The corrected table and figure legends are printed below

Table 5. Nonhematologic Adverse Events in 41 Patients Who Received Study Drug

	Placebo + Filgrastim $(n = 10)$	PEG-rHuMGDF* + Filgrastim (n = 31)
Local		
Injection site reaction	0 (0)†	2 (6)
Systemic		
Lethargy/drowsiness	1 (10)	17 (55)
Hot flushes/fever*	6 (60) [1]‡	8 (26)
Bone pain	3 (30)	8 (26)
Neurologic		
Headache	6 (60)	13 (42)
Dizziness	3 (30)	12 (38)
Respiratory		
Cough	1 (10)	8 (26)
Dypsnea	3 (30)	12 (38) [3]
Gastrointestinal		
Nausea and/or vomiting	9 (90) [1]	22 (71) [4]
Diarrhea	3 (30)	15 (48) [2]
Mucositis	5 (50)	8 (26)
Metabolic		
Hypokalemia	2 (20)	5 (16)
Thromboembolic		
Thrombophlebitis	0 (0)	1 (3)
Pulmonary embolism	0 (0)	1 (3) [1]§

There was no difference in the frequency of any toxicity between patients given placebo and those administered PEG-rHuMGDF (Fisher's Exact test).

Fig 1. Median platelet counts after chemotherapy for patients given PEF-rHuMGDF 0.03 ( $\blacktriangle$ , n = 3), 0.1  $\mu$ g/kg ( $\blacksquare$ , n = 3), 0.3  $\mu$ g/kg ( $\blacktriangledown$ , n = 3), 1.0  $\mu$ g/kg ( $\spadesuit$ , n = 11), 3.0  $\mu$ g/kg ( $\spadesuit$ , n = 7), or 5.0  $\mu$ g/kg ( $\square$ , n = 4), and placebo ( $\triangle$ , n = 10). The horizontal bar indicates a platelet count of 100  $\times$  10°/L.

Fig 2. (A) shows platelet recovery for all patients given PEG-rHuMGDF ( $\blacksquare$ , n = 25) and those given placebo ( $\triangle$ , n = 10). (B) shows the Kaplan-Meier plot of the probability of recovery to baseline platelet counts in these groups. The median time to baseline counts was 18 days for PEG-rHuMGDF and 22 days for placebo (P = .014, log rank test).

<sup>\*</sup>Not related to neutropenia.

<sup>†</sup>No. of patients (percentage).

<sup>‡</sup>Numbers in brackets represent number of events that were grade

<sup>3.</sup> 

<sup>§</sup>This event was grade 4.