

Active specific immunotherapy in patients with melanoma. A clinical trial with mouse antiidiotypic monoclonal antibodies elicited with syngeneic anti-high-molecular-weight-melanoma-associated antigen monoclonal antibodies.

J Clin Invest. 1991;**87**(2):757-757. <https://doi.org/10.1172/JCI114952C1>.

Correction

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A. Mittelman, Z. J. Chen, T. Kageshita, H. Yang, M. Yamada, P. Baskind, N. Goldberg, C. Puccio, T. Ahmed, Z. Arlin, and S. Ferrone.

The Journal of Clinical Investigation, Volume 86, No. 6, December 1990.

Page 2142.

Due to the publisher's error, a revised Figure 8 was not included in the article. It is printed below.

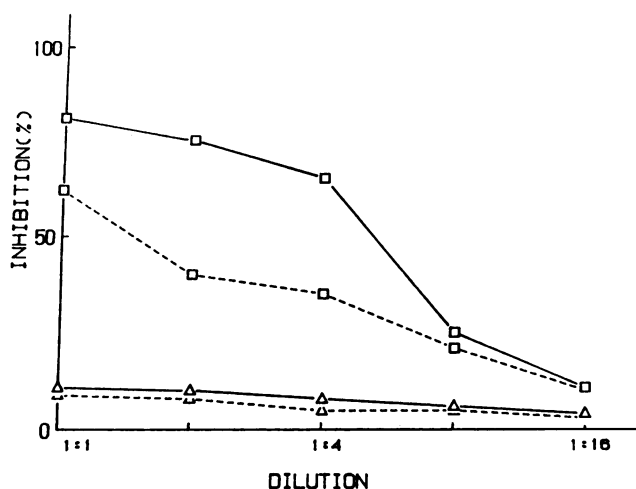


Figure 8. Inhibition of the binding of anti-HMW-MAA MAb 225.28 to cultured human melanoma cells Colo 38 by sera from patient MM immunized with the mouse antiidiotypic MAb MF11-30. Sera were obtained on day 42 (---) and 63 (—) from patient MM who had been immunized on day 0, 7, 28, and 42 with subcutaneous injections of MAb MF11-30 (2 mg per injection). Cultured melanoma cells Colo 38 were incubated with patient MM's serum for 2 h at 4°C. Then the supernatant was carefully removed and cells were added with ^{125}I -anti-HMW-MAA MAb 225.28 (□). Following an additional 2-h incubation at 4°C, cells were washed with PBS supplemented with 1% BSA (PBS-BSA) and bound radioactivity was measured in a gamma counter. Results are expressed as percent inhibition of the binding of ^{125}I -labeled MAb to melanoma cells as compared with binding performed in the presence of preimmune serum. ^{125}I -anti-ICAM-1 MAb CL207.14 (Δ) was used as a specificity control.