Table *IFNγ-secreting cells (s.c.) in multiple sclerosis patients treated with IFNβ-1b*

Group	Number of IFNγ s.c. cultured ex vivo in medium at baseline	Number of IFNγ s.c. cultured ex vivo in medium while on IFNβ-1b
MS patients at 24 hr	7.5 ± 2.17 (12)*	9.14 ± 3.06
MS patients at 1 wk	4.0 ± 0.97 (12)	6.29 ± 1.13
MS patients at 2–10 wk	2.76 ± 0.86 (9)	4.34 ± 1.32
MS patients beyond 3 mo	5.71 ± 1.97 (7)	1.85 ± 0.37
MS patients receiving prednisone 24 hr-3 mo	4.33 ± 1.27 (9)	3.14 ± 0.6

* Mean number of IFNγ s.c./10^5 cells ± S.E.M. Number of subjects in parentheses.

erable variations among the groups studied at baseline. Nonetheless, the number of cells secreting IFNγ "spontaneously" is higher 24 hours after starting IFNβ-1b treatment than for the same patients at baseline. The same holds at one week, and at 2–10 weeks. For patients studied one week after beginning treatment, the increase in cells that secrete IFNγ "spontaneously" is significant when compared to baseline values for the same patients ($p < 0.02$). IFNγ secreting cells were lower than baseline values in six of the seven patients studied after more than 3 months of treatment, but the difference failed to reach statistical significance. Prednisone-treated patients showed no change in IFNγ secreting cell numbers while on treatment, as was the case with ConA stimulation as well. Overall, the data complement and reinforce those presented in the paper.

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References

1. Dayal AS, Jensen MA, Lledo A, Arnason BGW. Interferon-gamma-secreting cells in multiple sclerosis patients treated with interferon beta-1b. Neurology 1995;45:2173–2177.
2. Farrar MA, Schreiber RD. The molecular cell biology of interferon-gamma and its receptor. Annu Rev Immunol 1993;11:571–611.
3. Panitch HS, Hirsch RL, Haley AS, Johnson KP. Exacerbations of multiple sclerosis in patients treated with gamma interferon. Lancet 1987; 1:893–895.
4. Olsson T, Zhi WW, Hojeberg B, Kostulas V, Jiang YP, Anderson G, Ekre HP, Link H. Autoreactive T lymphocytes in multiple sclerosis determined by antigen-induced secretion of interferon-gamma. J Clin Invest 1990;86:981–985.
5. Sun JB, Olsson T, Wang WZ, Xiao BG, Kostulas V, Fredrikson S, Ekre HP, Link H. Autoreactive T and B cells responding to myelin proteolipid protein in multiple sclerosis and controls. Eur J Immunol 1991;21: 1461–1468.
6. Sun J, Link H, Olsson T, Xiao BG, Andersson G, Ekre HP, Linington C, Diener P. T and B cell responses to myelin-oligodendrocyte glycoprotein in multiple sclerosis. J Immunol 1991;146:1490–1495.
7. Vartanian T, Li Y, Zhao MJ, Stefansson K. Interferon-gamma-induced oligodendrocyte cell death—implications for the pathogenesis of multiple sclerosis. Molecular Medicine 1995;1:732–743.
8. Mustafa MI, Diener P, Hojeberg B, Van der Meide P, Olsson T. T cell immunity and interferon-gamma secretion during experimental allergic encephalomyelitis in Lewis rats. J Neuroimmunol 1991;31:165–177.
9. Olsson T. Cytokine-producing cells in experimental autoimmune encephalomyelitis and multiple sclerosis. Neurology 1995;45:S11–S15.
10. Roosnek EE, Brouwer MC, Aarden LA. T cell triggering by lectins. I. Requirements for interleukin 2 production; lectin concentration determines the accessory cell dependency. Eur J Immunol 1985;15:652–656.
11. Weinstock-Guttman B, Ransohoff RM, Kinkel RP, Rudick RA. The interferons: biological effects, mechanisms of action, and use in multiple sclerosis. Ann Neurol 1995;37:7–15.

Corrections

In "Report of the AAN Task Force on access to health care: The effect of no personal health insurance on health care for people with neurologic disorders," that appeared in the May issue (Neurology 1996;47:1471–1480), the name of a member of the Task Force was inadvertently omitted from **Appendix 1.** The omitted individual is Edgar J. Kenton, III, MD, Division of Neurology, Lankenau Hospital, Philadelphia, PA.

The following is an addendum to the article, "Obsessive-compulsive disorder associated with brain lesions: Clinical phenomenology, cognitive function, and anatomic correlates," by Berthier et al., that appeared in the August issue (Neurology 1996;47:353–361): **Note.** Readers can obtain 8 pages of supplementary material from the National Auxiliary Publications Service, c/o Microfiche Publications, PO Box 3513, Grand Central Station, New York, NY 10163-3513. Request document no. 05331. Remit with your order (not under separate cover), in US funds only, $7.75 for photocopies or $5.00 for microfiche. Outside the United States and Canada, add postage of $4.50 for the first 20 pages and $1.00 for each 10 pages of material thereafter, or $1.75 for the first microfiche and $.50 for each fiche thereafter. *There is a $15.00 invoicing charge on all orders filled before payment.*