

[18] Electrophoretic Separation of Lymphoid Cells

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Introduction

The dissection of a complex system into its components, their separate analysis, and the controlled reconstitution is a scientific principle that has been used quite successfully in biochemistry. Its application in immunology calls for techniques for large-scale separation of highly pure and functionally intact lymphocyte subpopulations. Since most lymphocyte functions are mediated through their cell surface, a separation based on differences in cell membrane properties, such as surface charge, seems to be especially promising.

Preparative electrophoresis of cells became possible with the development of free-flow electrophoresis by K. Hannig.¹⁻³ The application of this method to the separation of lymphocytes is closely connected with the name of our colleague, the late K. Zeiller. He and a number of workers in other laboratories have demonstrated the potential of free-flow electrophoresis to efficiently separate murine T and B cells, as well as lymphocyte subpopulations at different stages of activation or differentiation (reviewed in refs. 4-9). Although most investigations have dealt with lymphocytes from mice and rats, lymphoid cells from humans,^{5,10-12} non-

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⁴ K. Zeiller, *Behring Inst. Mitt.* **52**, 11 (1972).

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⁶ G. V. Sherbet, ed., "The Biophysical Characterization of the Cell Surface," Chapter 4, p. 36. Academic Press, New York, 1978.

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¹⁰ G. Stein, H. D. Flad, R. Pabst, and F. Trepel, *Biomedicine* **19**, 388 (1973).

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