phagocytic activity even after a long in vitro culturing period of 2 wk. Our procedure will therefore be useful for investigating research problems involving macrophages.

## V. REFERENCES

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[1] K. C. Biological, Inc., Lenexa, KS.
[2] Flow Laboratories, Inglewood, CA.
[3] Abbott Laboratories, North Chicago, IL.
[4] McGaw Laboratories, Irving, CA.
[5] Sigma Chemical Co., St. Louis, CA.
[6] American Scientific Products, Denver, CO.
[7] Becton Dickinson and Company, Rutherford, NJ.
[8] Moore Push Pin Co., Wyndmoor, PA.
[9] Ethicon, Inc., Sommerville, NJ.
[10] American Sterilizer, Erie, PA.
[11] Forma Scientific, Inc., Marietta, OH.
[12] Dremel Electric Co., Racine, WI.
[13] Dupont Company, Instruments Products, Biomedical Division, Newton, CT.
[14] Inbred colony obtained from Inhalation Toxicology Research Institute, Albuquerque, N.M.

## ERRATUM

**Preparation of Primary Monolayer Cultures of Adult Rat Hepatocytes**
*G. L. Engelmann and J. A. Fierer*
Volume 7, No. 4, pages 169–173; 1982. (Procedure Number 41147). In III PROCEDURE, item A. 2. b. (page 170, column 2) should read:

b. To 225 ml add 140 mg $CaCl_2$ (Solution A).