DEPARTMENT OF HEALTH AND HUMAN SERVICES
WASHINGTON, D.C.

BIOLOGICS LICENSE
FOR THE MANUFACTURE OF
BIOLOGICAL PRODUCTS

This is to certify that Biologics License No. 249 is hereby issued to Northern Illinois Blood Bank, Inc., the manufacturer at Rockford, Illinois pursuant to Section 351 of the Public Health Service Act, approved July 1, 1944 (58 Stat. 702, 42 U.S.C. 262), as amended, and the regulations thereunder. The license authorizes the maintenance of an establishment(s) for the propagation or manufacture and preparation for sale, barter, or exchange in the District of Columbia, or for sending, carrying, or bringing for sale, barter, or exchange from any State or possession into any other State or possession or into any foreign country, or from any foreign country into any State or possession, any approved virus, therapeutic serum toxin, antitoxin, vaccine, blood, blood component or derivative, allergenic product, or analogous product, or arsphenamine or its derivatives, for which the licensee has demonstrated compliance to establishment and product standards.

Date September 1, 1998

[Signature]
Director, Center for Biologics Evaluation and Research
Food and Drug Administration