DEPARTMENT OF HEALTH AND HUMAN SERVICES
PUBLIC HEALTH SERVICE
FOOD AND DRUG ADMINISTRATION
Washington, D.C.

ESTABLISHMENT LICENSE
FOR THE MANUFACTURE OF
BIOLOGICAL PRODUCTS

This is to certify that Establishment License No. _______ 354 _______ is hereby issued
to ______________________, the manufacturer,
located at ______________________, through the establishment
identified as ______________________,
located at ______________________,
pursuant to Section 351 of the Public Health Service Act, approved July 1, 1944 (38 Stat. 702, 42 U.S.C. 262), as amended, and the
regulations thereunder. The license authorizes the manufacturer to maintain an establishment 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 virus, therapeutic serum, toxin, antitoxin, vaccine, blood, blood component or derivative, allergenic product, or analogous
product, or arsenical or its derivatives, for which the manufacturer holds an suspended and unrevoked product license issued by
the Secretary of Health and Human Services pursuant to said Act and regulations.

Date: July 29, 1997

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