

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

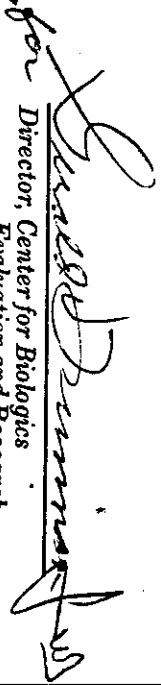
# ESTABLISHMENT LICENSE

FOR THE MANUFACTURE OF  
BIOLOGICAL PRODUCTS

This is to certify that Establishment License No. 688 is hereby issued  
to \_\_\_\_\_, the manufacturer,  
located at \_\_\_\_\_, Knoxville, Tennessee  
\_\_\_\_\_ through the establishment  
identified as \_\_\_\_\_, Medic, Inc.  
located at \_\_\_\_\_, Knoxville, Tennessee

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 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 arsenamine or its derivatives, for which the manufacturer holds an unsuspended and unrevoked product license issued by the Secretary of Health and Human Services pursuant to said Act and regulations.

Date June 6, 1989

  
for Director, Center for Biologics  
Evaluation and Research  
Food and Drug Administration

