

NRC Form 483  
1-76  
CFR 31

Approved by GAO  
38-R0160

U.S. NUCLEAR REGULATORY COMMISSION  
REGISTRATION CERTIFICATE—IN VITRO TESTING  
WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE

Section 31.11 of 10 CFR 31 establishes a general license authorizing physicians, clinical laboratories, and hospitals to possess certain small quantities of byproduct material for *in vitro* clinical or laboratory tests not involving the internal or external administration of the byproduct material or the radiation therefrom to human beings or animals. Possession of byproduct material under 10 CFR 31.11 is not authorized until the physician, clinical laboratory, or hospital has filed an application and received from the Commission a validated copy of NRC Form 483 with registration number.

Washington Square Clinic, P.C.  
1200 West Twelve Mile Road  
Madison Heights, MI 48071

... pursuant to  
... byproduct materials for  
... authorized to dispense

8777

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

APRIL 10, 1991

BRENDA E. BROWN

Date 1-14-91

Joseph A. Mabilia, D.O., President

WARNING—18 U.S.C., Section 1001: It is prohibited to knowingly and willfully provide false information to any agency or officer of any agency.