

Bankaci Medical Associates, Inc.

MURAT BANKACI, M.D., F.A.C.S.

OTOLARYNGOLOGY

HEAD AND NECK SURGERY

609 NORTH CHURCH STREET, MOUNT PLEASANT, PENNSYLVANIA 15666 547-4575

833 RUGH STREET, GREENSBURG, PENNSYLVANIA 15601 832-9665

February 23, 1987

U.S. Nuclear Regulatory Commission
631 Park Avenue
King of Prussia, PA 19406

To Whom it may concern:

Enclosed is a revised NRC Form 483 due to a change of address:

Previous Address was: Murat Bankaci, M.D.
110 South Arch Street
Suite 1A
Connellsville, PA 15425

New Address is: Murat Bankaci, M.D., F.A.C.S.
609 North Church Street
Mount Pleasant PA 15666

Registration number is 5852.

Thank you,

Murat Bankaci, M.D., F.A.C.S.

MB/trq

9104110169 910220
PDR FOIA
ASARCH91-38 PDR

9104110169

ADDRESS Change

Bankaci Medical Associates, Inc.

MURAT BANKACI, M.D., F.A.C.S.

OTOLARYNGOLOGY

HEAD AND NECK SURGERY

609 NORTH CHURCH STREET, MOUNT PLEASANT, PENNSYLVANIA 15666 547-4875
533 RUGH STREET, GREENSBURG, PENNSYLVANIA 15601 832-9688

FEBRUARY 5, 1987

TO WHOM IT MAY CONCERN:

I PRESENTLY POSSESS A REGISTRATION CERTIFICATE, IN VITRO TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE, N.C.R. FORM H83. I WOULD LIKE TO MAKE A CHANGE OF ADDRESS FOR YOUR RECORDS. MY REGISTRATION NUMBER IS: 5852. THE DATE OF ISSUE WAS MARCH 26, 1981. THE PROPER ADDRESS IS: MURAT BANKACI M.D.,
609 NORTH CHURCH STREET
MOUNT PLEASANT, PA 15666

THANK YOU,

Murat Bankaci M.D.
MURAT BANKACI M.D.
BANKACI MEDICAL ASSOC., INC.
609 NORTH CHURCH STREET
MOUNT PLEASANT, PA 15666

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 NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

Murat Benkeci, M.D.
110 S. Arch Street, Suite 1-A
Connellsville, PA 15425


3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct material for (please check one block only):
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.
4. To be completed by the Nuclear Regulatory Commission.

INSTRUCTIONS

1. Submit this form in triplicate to:
Office of Nuclear Material Safety and Safeguards
ATTN: Radioisotopes Licensing Branch
U.S. Nuclear Regulatory Commission
Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant: physician, clinical laboratory, or hospital for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. If NRC a registration number will be assigned and a validated copy of NRC Form 483 will be returned.

Registration number: **5852**

FOR THE U. S. NUCLEAR REGULATORY COMMISSION



Shirley A. Crutchfield, + March 26, 1981

(If this is an initial registration, leave this space blank - number to be assigned by NRC. If this is a change of information from a previous's registered general licensee, include your registration number.)

5. If place of use is different from address in Item 1, please give complete address:

6. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form) and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date 3/30/81

By *Shirley A. Crutchfield*
Signature of person filing form

Murat Benkeci, M.D.
Printed name and title of position of person filing form

WARNING--18 U.S.C., Section 1001, Act of June 25, 1948, 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

U.S. ATOMIC ENERGY 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 Form AEC-483 and received from the Commission a validated copy of Form AEC-483 with registration number.

INSTRUCTIONS

Submit this form in triplicate to: United States Atomic Energy Commission, Washington, D.C. 20545, Attention: Director, Division of Materials Licensing. A registration number will be assigned and a validated copy of Form AEC-483 will be returned.

1. Please print or type within the shaded area, below, the name and address (including ZIP Code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed.

Dr. Victor A. Bentley - Director #37-01873-01
Canaan Valley Memorial Hospital AEC Home
1044 Franklin Street
Johnstown, Penna. 15480


2. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one):

- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
- b. The above-named clinical laboratory.
- c. The above-named hospital.

3. To be completed by the Atomic Energy Commission

Registration number 1275

U. S. ATOMIC ENERGY COMMISSION



BY: *CA* **Clarence A. Hebron** **Sept. 24, 1971**
(Leave this space blank—number to be assigned by AEC.)

4. If place of use is different from address in Item 1, please give complete address:

5. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director, Division of Materials Licensing, within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date 9-21-71

By *Victor A Bentley*
Signature of person filing form

Victor A Bentley, Director
Printed name and title or position of person filing form

U.S. ATOMIC ENERGY 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 Form AEC-463 and received from the Commission a validated copy of Form AEC-463 with registration number.

INSTRUCTIONS

Submit this form in triplicate to: United States Atomic Energy Commission, Washington, D.C. 20545, Attention: Director, Division of Materials Licensing. A registration number will be assigned and a validated copy of Form AEC-463 will be returned.

Please print or type within the shaded area, below, the name and address (including ZIP Code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed.

Baseline Memorial Hospital
Oakhill Drive
Grove City, Pennsylvania 16127

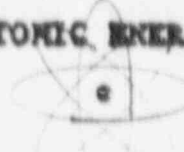
3. To be completed by the Atomic Energy Commission

2. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one):

- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 b. The above-named clinical laboratory.
 c. The above-named hospital.

Registration number: 0212

U. S. ATOMIC ENERGY COMMISSION



BY: John F. Schneider
(Print this name and number as validated by AEC)

4. If place of use is different from address in Item 1, please give complete address:

5. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director, Division of Materials Licensing, within 30 days from the effective date of such change.
d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form), and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date January 20, 1969

By Frederick J. Raisch
Signature of person filing form

Frederick J. Raisch, M.D. Pathologist, Director of Laboratories

Printed name and title or position of person filing form

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 Form AEC-483 and received from the Commission a validated copy of Form AEC-483 with registration number.

IRWIN BECKER, M.D.
216 WEST CASTLETON AVE.
PHILADELPHIA, PENN. 19144

3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct materials for (please check one block only)
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.
4. To be completed by the Atomic Energy Commission

INSTRUCTIONS

1. Submit this form in triplicate to:
Director of Licensing
ATTN: Materials Branch
Regulation
U.S. Atomic Energy Commission
Washington, D.C. 20545
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (At AEC, a registration number will be assigned and a validated copy of Form AEC-483 will be returned.)

Registration number:	2638
U. S. ATOMIC ENERGY COMMISSION	
	
BY: ^{CA} Clarence A. Hebron	6/6/74
<small>(If this is an initial registration, leave this space blank - number to be assigned by AEC. If this is a change of information from a previously registered general licensee, include your registration number.)</small>	

5. If place of use is different from address in Item 1, please give complete address:

6. Certification

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Licensing, within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date 5/17/74

By Irwin Becker, M.D.
Signature of person filing form

Printed name and title or position of person filing form

WARNING—18 U.S.C., Section 1001, Act of June 25, 1948; 62 Stat. 749; makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

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 NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

Albert M. Benshoff, M.D.
Mercy Hospital
1020 Franklin St.
Johnstown, Pa. 15905


- 3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct materials for *(please check one block only)*
 - a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.
- 4. To be completed by the Nuclear Regulatory Commission:

INSTRUCTIONS

- 1. Submit this form in triplicate to:
Office of Nuclear Material Safety and Safeguards
ATTN: Radioisotopes Licensing Branch
U.S. Nuclear Regulatory Commission
Washington, D.C. 20555
- 2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (A) NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.

Registration number: **5548**

FOR THE U. S. NUCLEAR REGULATORY COMMISSION



Shirley A. Crutchfield** **July 16, 1980**

(If this is an initial registration, leave this space blank -- number to be assigned by NRC. If this is a change of information from a previously registered general licensee, include your registration number.)

5. If place of use is different from address in Item 1, please give complete address:

6. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form), and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date 6/29/80

By Albert M. Benshoff, M.D.
Signature of person filing form.
Director, Clinical Laboratory
Mercy Hospital

Permit under: Department of Clinical Laboratories, Commonwealth of Pennsylvania, Dept. of Health

Printed name and title of person filing form

WARNING—18 U.S.C. Section 1001, Act of June 25, 1948, 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

U.S. ATOMIC ENERGY 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 Form AEC-483 and received from the Commission a validated copy of Form AEC-483 with registration number.

INSTRUCTIONS

Submit this form in triplicate to: United States Atomic Energy Commission, Washington, D.C. 20545, Attention: Director, Division of Materials Licensing. A registration number will be assigned and a validated copy of Form AEC-483 will be returned.

1. Please print or type within the shaded area, below, the name and address (including ZIP Code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed.


Edward J. Benz, M. D.
The Laboratory of Clinical Pathology, Inc.
19 East Elizabeth Avenue
Bethlehem, Pennsylvania 18018

3. To be completed by the Atomic Energy Commission

2. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one):
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.

Registration number: **1429**

U. S. ATOMIC ENERGY COMMISSION



BY: *CAH* **Clarence A. Habron** **Oct. 4, 1971**

(Leave this space blank—number to be assigned by AEC)

4. If place of use is different from address in Item 1, please give complete address

5. Certification:

- I hereby certify that:
- a. All information in this registration certificate is true and complete.
 - b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
 - c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director, Division of Materials Licensing, within 30 days from the effective date of such change.
 - d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form), and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date September 24, 1971

By

Edward J. Benz
Signature of person filing form

Edward J. Benz, M. D.

Printed name and title or position of person filing form

WARNING—18 U.S.C. Section 1001, Act of June 25, 1948, 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

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, hospitals, and veterinarians in the practice of veterinary medicine 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, hospital, or veterinarian in the practice of veterinary medicine, has filed NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

Dr. Alan S. Berger
525 Jamestown Avenue
Suite 104
Philadelphia, PA 19128


3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct materials for (please check one block only)
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.
 - d. Veterinarian in the practice of veterinary medicine.
4. To be completed by the Nuclear Regulatory Commission.

INSTRUCTIONS

1. Submit this form in triplicate to:
Office of Nuclear Material Safety and Safeguards
ATTN: Material Licensing Branch
U.S. Nuclear Regulatory Commission
Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, hospital, or veterinarian in the practice of veterinary medicine for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (At NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.)

Registration number: 8505

FOR THE U.S. NUCLEAR REGULATORY COMMISSION



(If this is an initial registration, leave this space blank. If number to be assigned by NRC. If this is a change of information from a previously registered general license, include your registration number.)

5. If place of use is different from address in Item 1, please give complete address:

6. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct material.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date X 2-13-86

By Alan S. Berger

Printed name and title or position of person filing form

WARNING— 18 U.S.C., Section 1001; Act of June 25, 1948; 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

REGISTRATION CERTIFICATE--IN VITRO TESTING
WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE

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Whenever the words "Atomic Energy Commission" or "Commission" appear in this registration, they mean the Nuclear Regulatory Commission created by Public Law 93-438 and Executive Order No. 11834.

Berks Clinical Laboratories, Inc.
107 S. 5th St.
Reading, PA. 19602

3. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one block only)

- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
- b. The above-named clinical laboratory.
- c. The above-named hospital.

4. To be completed by the Atomic Energy Commission

INSTRUCTIONS

1. Submit this form in triplicate to:
United States Atomic Energy Commission
Attention: Directorate of Licensing,
Materials Branch
Washington, D.C. 20545
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (At AEC, a registration number will be assigned and a validated copy of Form AEC-483 will be returned.)

Registration number: **3986**

For the U. S. Nuclear Regulatory Commission

Shirley A. Windley
Shirley A. Windley

March 7, 1977
Leave this space blank - number to be assigned by AEC.

5. If place of use is different from address in item 1, please give complete address:

Berks Clinical Laboratories, Inc.
c/o Cannon Laboratories, Inc.
Reading Municipal Airport
Reading, Pa. 19602

6. Certification:

I hereby certify that:

- a. All information on this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Directorate of Licensing, Materials Branch, within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form) and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date 3/1/77

By *Norman E. Kraus*
Signature of person filing form.

Norman E. Kraus - Director

Printed name and title of person filing form.

WARNING-18 U.S.C. Section 1001, Act of June 25, 1948 (62 Stat. 749) makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

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 NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

Howard J. Berman, M.D.
Director, Laboratory
Jeannette Dist. Memorial Hospital
600 Jefferson Ave.
Jeannette, Pa. 15644


3. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one block only)
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine
 - b. The above-named clinical laboratory
 - c. The above-named hospital
4. To be completed by the Nuclear Regulatory Commission

INSTRUCTIONS

1. Submit this form in triplicate to:
Office of Nuclear Material Safety and Safeguards
ATTN: Radioisotopes Licensing Branch
U.S. Nuclear Regulatory Commission
Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (At NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.)

Registr. number: 6780

FOR THE U. S. NUCLEAR REGULATORY COMMISSION



Shirley A. Crutchfield November 17, 1983
(If this is an initial registration, leave this space blank — number to be assigned by NRC. If this is a change of information from a previously registered general licensee, include your registration number.)

5. If place of use is different from address in item 1, please give complete address:

6. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form), and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date 11/7/83

By Howard J. Berman
Signature of person filing form

Howard J. Berman M.D., Director, Laboratory

Printed name and title of person filing form

WARNING—18 U.S.C., Section 1001, Act of June 25, 1948, 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

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, hospitals, and veterinarians in the practice of veterinary medicine 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, hospital, or veterinarian in the practice of veterinary medicine, has filed NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

**Berwick Hospital Laboratory
The Berwick Hospital
701 E. 10th Street
Berwick, Penna. 18603**

3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct materials for (please check one block only)


- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.
 - d. Veterinarian in the practice of veterinary medicine.
4. To be completed by the Nuclear Regulatory Commission.

INSTRUCTIONS

1. Submit this form in triplicate to:
Office of Nuclear Material Safety and Safeguards
ATTN: Material Licensing Branch
U.S. Nuclear Regulatory Commission
Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant: physician, clinical laboratory, hospital, or veterinarian in the practice of veterinary medicine for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (At NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.

Registration number: **6337**

FOR THE U. S. NUCLEAR REGULATORY COMMISSION



Shirley A. Grutchfield May 6, 1982

If this is an initial registration, leave this space blank. If this is a change of information from a previously registered general license, include your registration number.

5. If place of use is different from address in Item 1, please give complete address:

Same As Above

6. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct material.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date **5/3/82**

By **Robert G. Hunter MD**

Robert G. Hunter, M. D., Chief of Pathology

Printed name and title or position of person filing form

WARNING— 18 U.S.C., Section 1001, Act of June 25, 1948; 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

U.S. ATOMIC ENERGY 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 Form AEC-483 and received from the Commission a validated copy of Form AEC-483 with registration number.

INSTRUCTIONS

Submit this form in triplicate to: United States Atomic Energy Commission, Washington, D.C. 20545, Attention: Director, Division of Materials Licensing. A registration number will be assigned and a validated copy of Form AEC-483 will be returned.

1. Please print or type within the shaded area, below, the name and address (including ZIP Code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed.

Richard P. Bandle, M. D.
Pathological Department
The Pottsville Hospital and Warne Clinic
Haugh Church & Jackson Sts.
Pottsville, Pa. 17854

Wherever the words "Atomic Energy Commission" or "Commission" appear in this registration, they mean the Nuclear Regulatory Commission created by Public Law 93-433 and Executive Order No. 11834.

5. To be completed by the Atomic Energy Commission

2. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one):

- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine
- b. The above-named clinical laboratory.

The above-named hospital

Registration number: 3462
For the U. S. Nuclear Regulatory Commission
EY: Clarence A. Hebron 12/12/75
(Leave this space blank—number to be assigned by AEC)

4. If place of use is different from address in Item 1, please give complete address.

1. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
- c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director, Division of Materials Licensing, within 30 days from the effective date of such change.
- d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date: November 24, 1975

By: *Richard P. Bandle*
Signature of person filing form

Richard P. Bandle, M. D.

Printed name and title in position of person filing form

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, hospitals, and veterinarians in the practice of veterinary medicine 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, hospital, or veterinarian in the practice of veterinary medicine, has filed NRC Form 483 and received from the Commission a validated copy of NRC Form 483 with registration number.

BIODECISION LABORATORIES
5900 PENN AVENUE
PITTSBURGH, PA 15206

3. I hereby apply for a registration number pursuant to §31.11, 10 CFR 31 for use of byproduct materials for (please check one block only)
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.
 - d. Veterinarian in the practice of veterinary medicine.
4. To be completed by the Nuclear Regulatory Commission.

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ATTN: Material Licensing Branch
U.S. Nuclear Regulatory Commission
Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, hospital, or veterinarian in the practice of veterinary medicine for whom or for which this registration form is filed. Position the first letter of the address below the left dot and do not extend the address beyond the right dot. (A: NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.)

Registration number: 8180

U.S. NUCLEAR REGULATORY COMMISSION



(If this is an initial registration, leave this space blank — number to be assigned by NRC. If this is a change of information from a previously registered general license, include your registration number.)

5. If place of use is different from address in Item 1, please give complete address:

6. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
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- d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date 3.24.87

By A. Bacharach

ALFRED BACHARACH, Ph.D., DIRECTOR OF RESEARCH AND DEVELOPMENT

Printed name and title of person filing form

WARNING— 18 U.S.C., Section 1001: Act of June 25, 1948, 62 Stat. 749, makes it a criminal offense to make a willfully false statement or representation to any department or agency of the United States as to any matter within its jurisdiction.

U.S. ATOMIC ENERGY COMMISSION
**REGISTRATION CERTIFICATE—IN VITRO TESTING
WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE**

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INSTRUCTIONS

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1. Please print or type within the shaded area, below, the name and address (including ZIP Code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed.

Medicinal Laboratories, Inc.
~~1251 Water Road~~ 491 Allendale Road
~~211 ...~~, ~~Dallas~~, 49141
King of Prussia, Pennsylvania 19406

*Name change from
Bernard L. Richmond to
George A. Adams as
per telephone call 8/1/72
E. Hahn*

2. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (please check one):
- a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
 - b. The above-named clinical laboratory.
 - c. The above-named hospital.

3. To be completed by the Atomic Energy Commission

Registration number: 0383

U.S. ATOMIC ENERGY COMMISSION

BY: *John V. Schneider*
BY: **John V. Schneider**
(Leave this space blank—number to be assigned by AEC)

If place of use is different from address in Item 1, please give complete address:

5. Certification:

I hereby certify that:

- a. All information in this registration certificate is true and complete.
- b. The registrant has appropriate radiation measuring instruments to carry out the tests for which byproduct material will be used under the general license of 10 CFR 31.11. The tests will be performed only by personnel competent in the use of the instruments and in the handling of the byproduct materials.
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- d. I have read and understand the provisions of Section 31.11 of AEC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with those provisions as to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Atomic Energy Commission.

Date: August 21, 1972

By: *George A. Adams*
Signature of person filing form

George A. Adams
Director, Division of Materials Licensing

Printed name and title or position of person filing form

1-2-2000

JUL 21 1970

Biomedical Laboratories, Inc.
491 Allendale Road
King of Prussia, Pa. 19406

Attention: Mr. Bernard I. Diamond
President

Gentlemen:

Thank you for your letter dated July 11, 1970, regarding change in address.

Our records relating to your Registration Number 0383 for In Vitro Testing with Byproduct Material under General License have been changed accordingly.

Section 31.11 of 10 CFR 31 does not require issuance of a new registration number for address changes; therefore, it will not be necessary for you to return your current registration.

Sincerely yours,

Frank V. Malone, Chief
Administrative Services
Office of the Director
of Regulation

ADMEY	DE	ADM:DI
HSmith	CEdwards	FMalone
7/1/70	7/1/70	7/1/70

JUL 22 1970

Biomedical Laboratories, Inc.
491 Allendale Road
King of Prussia, Pa. 19406

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President

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Sincerely yours,

Frank W. Malone, Chief
Administrative Services
Office of the Director
of Regulation

OFFICE ▶	ADM:DR	DML	ADM:DR		
SURNAME ▶	HSmith:ac	CEdwards	FMalone		
DATE ▶	7/20/70	7/21/70	7/ /70		