

BYPRODUCT MATERIAL LICENSE
(Medical - Groups I & II)

Amendment No. 36

Pursuant to the Atomic Energy Act of 1954 and Title 10, Code of Federal Regulations, Chapter 1, Parts 30, 32, 33, 34, and 35, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, own, possess, transfer and import byproduct material listed below; and to use such byproduct material for the purpose(s) and at the place(s) designated below. This license shall be deemed to contain the conditions specified in Section 183 of the Atomic Energy Act of 1954 as amended, and is subject to all applicable rules, regulations, and orders of the Atomic Energy Commission now or hereafter in effect and to any conditions specified below.

Licensee		In accordance with application dated March 12, 1975,	
1. Department of the Army Fitzsimons Army Medical Center and U.S. Army Medical Research and Nutrition Laboratory Denver, Colorado 80240		3. License Number	05-00046-13 is amended in its entirety to read as follows:
		4. Expiration date	April 30, 1979
		5. Reference No.	
6. Byproduct material (element and mass number)	7. Chemical and/or physical form	8. Maximum amount of radioac- tivity which licensee may possess at any one time	
A. Any byproduct material listed in Groups I and II of Schedule A, Section 35.100 of 10 CFR 35	A. Any radio- pharmaceutical listed in Groups I and II of Schedule A, Section 35.100 of 10 CFR 35	A. As necessary for uses authorized in Subitem 9. A.	
B. Any byproduct material listed in Group III of Schedule A, Section 35.100 of 10 CFR 35	B. Any form listed in Group III of Schedule A, Section 35.100 of 10 CFR 35	B. 2 curies of each byproduct material authorized in Subitem 6.B.	
C. Any byproduct material listed in Group IV of Schedule A, Section 35.100 of 10 CFR 35	C. Any radio- pharmaceutical listed in Group IV of Schedule A, Section 35.100 of 10 CFR 35	C. As necessary for uses authorized in Subitem 9.C.	

Conditions numbered 1. printed on the reverse side of this page shall apply to this license.

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U. S. ATOMIC ENERGY COMMISSION
BYPRODUCT MATERIAL LICENSE

Supplementary Sheet

Continued from Page 1

05-00046-13

License Number _____

Amendment No. 36

Byproduct material
(element and mass number)

7. Chemical and/or physical form

8. Maximum amount of radio activity which
licensee may possess at any one timeD. Any byproduct
material listed
in Group V
of Schedule A,
Section 35.100
of 10 CFR 35E. Any byproduct
material listed
in Group VI of
Schedule A,
Section 35.100
of 10 CFR 35

F. Iodine 131

G. Iodine 125

H. Cesium 137

I. Any byproduct
material with
Atomic Nos. 1-83,
inclusive

J. Xenon 133

K. Hydrogen 3

L. Sodium 24

D. Any radio-
pharmaceutical
listed in Group
V of Schedule A,
Section 35.100
of 10 CFR 35E. Any sealed
source listed
in Group VI
of Schedule A,
Section 35.100
of 10 CFR 35

F. Thyroxine

G. Thyroxine

H. Any

I. Any

J. Free gas or in saline

K. Water

L. Sodium chloride

D. As necessary
for uses
authorized in
Subitem 9.D.E. 1 curie total
for all sources
authorized in
Subitem 6.E.

F. 2 millicuries

G. 1 millicurie

H. 1 millicurie

I. 500 millicuries each,
except: Hydrogen 3 -
5 curies. Total not to
exceed 10 curies

J. 2 curies

K. 25 millicuries

L. 1 millicurie

9. Authorized use

- A. Any diagnostic procedure listed in Groups I and II of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.
- B. Preparation and use of radiopharmaceuticals for any diagnostic procedure listed in Group III of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.
- C. Any therapeutic procedure listed in Group IV of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.
- D. Any therapeutic procedure listed in Group V of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.
- E. Any procedure listed in Group VI of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.

MATERIALS LICENSE

Supplementary Sheet

License Number 05-00046-13Docket or
Reference No. _____

Amendment No. 36

9. Authorized use (continued)

- F. and G. Determination of thyroxine turnover.
- H. Standard for assay of molybdenum content of eluate of molybdenum generator.
- I. Laboratory research in vitro and in lower animals, in vitro testing.
- J. Pulmonary function studies. Blood flow studies.
- K. Determination of total body water.
- L. Determination of total exchangeable sodium.

CONDITIONS

Wherever the words "Atomic Energy Commission" or "Commission" appear in this license, except where the context of their use refers to a fact or event prior to January 19, 1975, they mean the Nuclear Regulatory Commission created by Public Law 93-438 and Executive Order No. 11834.

- 10. The licensee shall comply with the provisions of Title 10, Chapter 1, Code of Federal Regulations, Part 19, "Notices, Instructions and Reports to Workers; Inspections" and Part 20, "Standards for Protection Against Radiation."
- 11. A. Byproduct material shall be used by, or under the supervision of, individuals designated by the Fitzsimons Army Medical Center Radioisotope Committee.

B. The use of byproduct material in or on humans shall be by a physician.
- 12. Byproduct material shall be used in accordance with the provisions of Section 35.14(b)(c)(e) and (f) of Title 10, Code of Federal Regulations, Part 35.
- 13. Except as specifically provided otherwise by this license, the licensee shall possess and use byproduct material described in Items 6, 7, and 8 of this license in accordance with statements, representations, and procedures contained in application dated February 22, 1974, and letters dated April 15, 1974 and July 30, 1975.

Date AUG 25 1975

For the U. S. Nuclear Regulatory Commission

Signed and Stamped By

John E. B. 201/201

by Materials BranchDivision of Materials and Fuel Cycle
Facility Licensing
Washington, D. C. 20555

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MATERIALS DATA INPUT

FILE COPY

A. TYPE OF ACTION AND IDENTIFICATION CODES

<input type="checkbox"/> NEW LICENSE	<input type="checkbox"/> AMENDMENT TO RENEW LICENSE	<input type="checkbox"/> AMENDMENT TO TERMINATE	<input type="checkbox"/> VOID	DOCKET NUMBER 030-01233	MAIL CONTROL NUMBER 55580	CHANGE NAME/ADDRESS <input type="checkbox"/>
<input type="checkbox"/> NEW LICENSE AND NEW LICENSEE	<input checked="" type="checkbox"/> OTHER AMENDMENT	<input type="checkbox"/> CLERICAL CHANGE NO AMENDMENT				

B. INDICATIVE INFORMATION:

INDIVIDUAL LICENSEE'S	NAME (LAST, FIRST, MIDDLE)	NAME (LAST, FIRST, MIDDLE)		
	NAME (LAST, FIRST, MIDDLE)	NAME (LAST, FIRST, MIDDLE)		
	NAME (LAST, FIRST, MIDDLE)	NAME (LAST, FIRST, MIDDLE)		
ORGANIZATION NAME (ALPHABETIC SEQUENCE)	Army, Department of the			
	DEPARTMENT OR BUREAU Fitzsimons Army Medical and U.S. Army Medical Research			
ADDRESS	BUILDING, STREET	CITY Denver	STATE KIBCO	ZIP CODE 80240
	TYPE OF APPLICANT <input type="checkbox"/> U.S. GOVERNMENT AGENCY <input type="checkbox"/> INDIVIDUAL LICENSEE <input checked="" type="checkbox"/> ORGANIZATIONAL LICENSEE	DATE REQUEST RECEIVED 04/02/75	INSTITUTION CODE 00046	PENDING PROG. CODE ACTUAL PROG. CODE
SECONDARY PROGRAM CODES AS REQUIRED:				
#1	#2	#3	#4	#5
LICENSE NUMBER 05-00046-13		DATE LICENSE ISSUED OR ACTION COMPLETED 08 25 75	EXPIRATION DATE 04-30-79	

C. STATISTICAL INFORMATION:

MEDICAL CATEGORY:					
<input type="checkbox"/> FOR HUMAN USE ONLY	<input checked="" type="checkbox"/> FOR HUMAN AND NONHUMAN USE	<input type="checkbox"/> FOR NONHUMAN USE ONLY			
POSSESSION OF THE MATERIAL IS AUTHORIZED IN ONE OF THE FOLLOWING AREAS:					
<input checked="" type="checkbox"/> SAME AS "STATE" IN ADDRESS	<input type="checkbox"/> ALL STATES	<input type="checkbox"/> ALL NON-AGREEMENT STATES			
AND/OR IN THE STATE(S), TERRITORY(S), COUNTRY CHECKED BELOW:					
ALABAMA -AL	GEORGIA -GA	MARYLAND -MD	NEW JERSEY -NJ	SOUTH CAROLINA -SC	WYOMING -WY
ALASKA -AK	HAWAII -HI	MASSACHUSETTS -MA	NEW MEXICO -NM	SOUTH DAKOTA -SD	
ARIZONA -AZ	IDAHO -ID	MICHIGAN -MI	NEW YORK -NY	TENNESSEE -TN	AMERICAN SAMOA -AS
ARKANSAS -AR	ILLINOIS -IL	MINNESOTA -MN	NORTH CAROLINA -NC	TEXAS -TX	CANAL ZONE -CZ
CALIFORNIA -CA	INDIANA -IN	MISSISSIPPI -MS	NORTH DAKOTA -ND	UTAH -UT	GUAM -GU
COLORADO -CO	IOWA -IA	MISSOURI -MO	OHIO -OH	VERMONT -VT	PUERTO RICO -PR
CONNECTICUT -CT	KANSAS -KS	MONTANA -MT	OKLAHOMA -OK	VIRGINIA -VA	VIRGIN ISLANDS -VI
DELAWARE -DE	KENTUCKY -KY	NEBRASKA -NB	OREGON -OR	WASHINGTON -WA	
WASHINGTON, DC -DC	LOUISIANA -LA	NEVADA -NV	PENNSYLVANIA -PA	WEST VIRGINIA -WV	CANADA -CH
FLORIDA -FL	MAINE -ME	NEW HAMPSHIRE -NH	RHODE ISLAND -RI	WISCONSIN -WI	

D. POSSESSION LIMITS OF SOURCE AND SPECIAL NUCLEAR MATERIALS AND TRITIUM

SOURCE MATERIAL CEILING		SNM CEILING						
<input type="checkbox"/> GRAMS	<input type="checkbox"/> KILOGRAMS	<input type="checkbox"/> GRAMS	<input type="checkbox"/> KILOGRAMS	<input type="checkbox"/> "X" HERE IF FOR POWER REACTOR				
AMOUNT	UNIT	CONFIG.	ENRICH.	MAT.	AMOUNT	UNIT	CONFIG.	ENRICH.
U5	<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS				<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS	
U3	<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS				<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS	
PU	<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS				<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS	
UR	<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS				<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS	
TH	<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS				<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS	
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	<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS				<input type="checkbox"/> G <input type="checkbox"/> kg	<input type="checkbox"/> S <input type="checkbox"/> UNS	
H3		<input type="checkbox"/> CURIES <input type="checkbox"/> MILLICURIES		RIS CODES				

MATERIALS DATA INPUT MEDICAL

5 MEDICAL BYPRODUCT
REFERENCE COPY

A. TYPE OF ACTION AND IDENTIFICATION CODES

<input type="checkbox"/> NEW LICENSE	<input type="checkbox"/> AMENDMENT TO RENEW LICENSE	<input type="checkbox"/> AMENDMENT TO TERMINATE	<input type="checkbox"/> VOID	DOCKET NUMBER 030-01233	MAIL CONTROL NUMBER 33708	CHANGE NAME ADDRESS <input type="checkbox"/>
<input type="checkbox"/> NEW LICENSE AND RENEW LICENSE	<input type="checkbox"/> OTHER AMENDMENT	<input type="checkbox"/> CLERICAL CHANGE NO AMENDMENT				

B. INDICATIVE INFORMATION

NAME (LAST, FIRST, MIDDLE)	NAME (LAST, FIRST, MIDDLE)			
NAME (LAST, FIRST, MIDDLE)	NAME (LAST, FIRST, MIDDLE)			
NAME (LAST, FIRST, MIDDLE)	NAME (LAST, FIRST, MIDDLE)			
ORGANIZATION NAME (ALPHABETIC SEQUENCE) Atomic Energy Commission of the U.S.				
DEPARTMENT OR BUREAU Department of Energy Medical and U.S. Army Medical Research				
ADDRESS	CITY	STATE	ZIP CODE	
Washington, D.C.	Washington	D.C.	20540	
TYPE OF APPLICANT <input type="checkbox"/> U.S. GOVERNMENT AGENCY <input type="checkbox"/> INDIVIDUAL LICENSEE <input checked="" type="checkbox"/> ORGANIZATIONAL LICENSEE	DATE REQUEST RECEIVED 05/20/75	INSTITUTION CODE 00001	PENDING PROC. CODE	ACTUAL PROC. CODE

C. SECONDARY PROGRAM CODES AS REQUIRED

NO.	PS	PS	PS
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FORM NUMBER	DATE LICENSE ISSUED OR ACTION COMPLETED	EXPIRATION DATE
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TYPE OF STUDY	FORM	USE	POSS. LIMIT
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10-00	IN VITRO STUDIES	Copy D, E, W, Y, N, L, M
10-01	IN VITRO STUDIES	
10-02	IN VITRO STUDIES	
10-03	IN VITRO STUDIES	
10-04	IN VITRO STUDIES	
10-05	IN VITRO STUDIES	
10-06	IN VITRO STUDIES	
10-07	IN VITRO STUDIES	
10-08	IN VITRO STUDIES	
10-09	IN VITRO STUDIES	
10-10	IN VITRO STUDIES	
10-11	IN VITRO STUDIES	
10-12	IN VITRO STUDIES	
10-13	IN VITRO STUDIES	
10-14	IN VITRO STUDIES	
10-15	IN VITRO STUDIES	
10-16	IN VITRO STUDIES	
10-17	IN VITRO STUDIES	
10-18	IN VITRO STUDIES	
10-19	IN VITRO STUDIES	
10-20	IN VITRO STUDIES	
10-21	IN VITRO STUDIES	
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10-97	IN VITRO STUDIES	
10-98	IN VITRO STUDIES	
10-99	IN VITRO STUDIES	
10-100	IN VITRO STUDIES	

- 9 L Determination of total body water
9 M Determination of total exchangeable sodium.
-

conditions

16. Fitzsimons Army Medical Center,
12101 East Colfax Ave, Aurora, Colorado

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9A. FAMC Radioisotope committee,

9B

18

12(h)(1) 12(d) Licensee

21 AA B

22

The licensee shall not use licensed material
in field applications except as provided
otherwise by specific conditions of this
License.

26

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63

16. appl. dated March 6, 1979
letter dated June 3, 1980

NRC FORM 754A
(5-78)

U.S. NUCLEAR REGULATORY COMMISSION

APPLICATION DATED

March 6, 1979

LETTER DATED

MEDICAL LICENSE DATA

RADIOACTIVE MATERIAL LISTED IN:		MARK ITEMS DESIRED "X"	MAXIMUM POSSESSION LIMITS (in millicuries)
1	10 CFR 35.100, SCHEDULE A, GROUP I	<input checked="" type="checkbox"/> A	As Needed
2	10 CFR 35.100, SCHEDULE A, GROUP II		As Needed
3	10 CFR 35.100, SCHEDULE A, GROUP III	<input checked="" type="checkbox"/> B	5 ci
4	10 CFR 35.100, SCHEDULE A, GROUP IV	<input checked="" type="checkbox"/> C	As Needed
5	10 CFR 35.100, SCHEDULE A, GROUP V	<input checked="" type="checkbox"/> D	As Needed
6	10 CFR 35.100, SCHEDULE A, GROUP VI	<input checked="" type="checkbox"/> E	2 ci
7	10 CFR 31.11 FOR IN-VITRO STUDIES		14 MCI
8	AMERICIUM - 241 ANATOMICAL MARKER		2 ci
9	XENON-133: As gas or gas in saline for blood flow and pulmonary function studies	<input checked="" type="checkbox"/> F	
10	IODINE-131: As iodide for treatment of hyperthyroidism and cardiac dysfunction		
11	IODINE-131: As iodide for treatment of thyroid carcinoma		
12	PHOSPHORUS-32: As soluble phosphate for treatment of polycythemia vera, leukemia and bone metastases		
13	PHOSPHORUS-32: As colloidal chromic phosphate for intracavitary treatment of malignant effusions		
14	GOLD-198: As colloid for intracavitary treatment of malignant effusions		
15	DEPLETED URANIUM FOR SHIELDING		

SYMBOL	CHEMICAL OR PHYSICAL FORM	POSSESSION LIMIT
G-3-83	standard tape G. Any	G. 500 mci, each
H-3	H. Any	H. 5 ci
I-Cs-137	I. Any	I. 1 mci
J-I-131	J. Thyroxine	J. 2 mci
K-I-125	K. Thyroxine	K. 1 mci
L-H-3	L. water	L. 25 mci
M-Na-24	M. NaCl	M. 1 mci

9 A-F. use standard medical uses

9 G-I RAD standard tape

9 J and K determination of thyroxine
Turnover

OVER

MAIL TO:

Arm V

DATE MAILED

BY

MM62

DATE COMPLETED

1-24-80

FROM

FITZSIMONS ARMY MEDICAL CENTER
DENVER, COLORADO

DATE OF DOCUMENT

DATE RECEIVED

12/8/76

NO

0690

LTR

MEMO

REPORT

OTHER

X

ORIG

CC

OTHER

X

ACTION NECESSARY

☒

CONCURRENCE

☐

DATE ANSWERED

NO ACTION NECESSARY

☐

COMMENT

☐

BY

CLASSIF

POST OFFICE

U

REG NO

FILE CODE

DESCRIPTION (Must Be Unclassified)

RESPONSE TO ENFORCMENT LTR DTD 11/24/
76 & INSP. HELD 11/11/76

REFERRED TO

DATE

RECEIVED BY

DATE

HOWARD

emb 12/8

BROWN

ENCLOSURES

D/F RELATING TO SECURITY OF RADIATION
AREA (HOT LAB)

VETTER (HAS COPY)

Spind

REMARKS

U. S. NUCLEAR REGULATORY COMMISSION

MAIL CONTROL FORM

FORM NRC 326
(1-75)

A/11/3