

APPLICATION FOR MATERIAL LICENSE

INSTRUCTIONS: SEE THE APPROPRIATE LICENSE APPLICATION GUIDE FOR DETAILED INSTRUCTIONS FOR COMPLETING APPLICATION. SEND TWO COPIES OF THE ENTIRE COMPLETED APPLICATION TO THE NRC OFFICE SPECIFIED BELOW.

APPLICATIONS FOR DISTRIBUTION OF EXEMPT PRODUCTS FILE APPLICATIONS WITH:

U.S. NUCLEAR REGULATORY COMMISSION
DIVISION OF FUEL CYCLE AND MATERIAL SAFETY, NMSS
WASHINGTON, DC 20546

ALL OTHER PERSONS FILE APPLICATIONS AS FOLLOWS, IF YOU ARE LOCATED IN:

CONNECTICUT, DELAWARE, DISTRICT OF COLUMBIA, MAINE, MARYLAND, MASSACHUSETTS, NEW HAMPSHIRE, NEW JERSEY, NEW YORK, PENNSYLVANIA, RHODE ISLAND, OR VERMONT, SEND APPLICATIONS TO:

U.S. NUCLEAR REGULATORY COMMISSION, REGION I
NUCLEAR MATERIALS SAFETY SECTION B
801 PARK AVENUE
KING OF PRUSSIA, PA 19406

ALABAMA, FLORIDA, GEORGIA, KENTUCKY, MISSISSIPPI, NORTH CAROLINA, PUERTO RICO, SOUTH CAROLINA, TENNESSEE, VIRGINIA, VIRGIN ISLANDS, OR WEST VIRGINIA, SEND APPLICATIONS TO:

U.S. NUCLEAR REGULATORY COMMISSION, REGION II
NUCLEAR MATERIALS SAFETY SECTION
101 MARIETTA STREET, SUITE 2800
ATLANTA, GA 30333

IF YOU ARE LOCATED IN:

ILLINOIS, INDIANA, IOWA, MICHIGAN, MINNESOTA, MISSOURI, OHIO, OR WISCONSIN, SEND APPLICATIONS TO:

U.S. NUCLEAR REGULATORY COMMISSION, REGION III
MATERIALS LICENSING SECTION
790 ROOSEVELT ROAD
GLEN ELLYN, IL 60137

ARKANSAS, COLORADO, IDAHO, KANSAS, LOUISIANA, MONTANA, NEBRASKA, NEW MEXICO, NORTH DAKOTA, OKLAHOMA, SOUTH DAKOTA, TEXAS, UTAH, OR WYOMING, SEND APPLICATIONS TO:

U.S. NUCLEAR REGULATORY COMMISSION, REGION IV
MATERIAL RADIATION PROTECTION SECTION
811 RYAN PLAZA DRIVE, SUITE 1000
ARLINGTON, TX 76011

ALASKA, ARIZONA, CALIFORNIA, HAWAII, NEVADA, OREGON, WASHINGTON, AND U.S. TERRITORIES AND POSSESSIONS IN THE PACIFIC, SEND APPLICATIONS TO:

U.S. NUCLEAR REGULATORY COMMISSION, REGION V
NUCLEAR MATERIALS SAFETY SECTION
1460 MARIA LAKE, SUITE 210
WALNUT CREEK, CA 94606

PERSONS LOCATED IN AGREEMENT STATES SEND APPLICATIONS TO THE U.S. NUCLEAR REGULATORY COMMISSION ONLY IF THEY WISH TO POSSESS AND USE LICENSED MATERIAL IN STATES SUBJECT TO U.S. NUCLEAR REGULATORY COMMISSION JURISDICTION.

1. THIS IS AN APPLICATION FOR (Check appropriate item.)

- A. NEW LICENSE
- B. AMENDMENT TO LICENSE NUMBER 24-24518-01
- C. RENEWAL OF LICENSE NUMBER _____

2. NAME AND MAILING ADDRESS OF APPLICANT (Include Zip Code)

St. Louis Regional Medical Center
5535 Delmar Blvd.
St. Louis, Missouri 63112

3. ADDRESS(ES) WHERE LICENSED MATERIAL WILL BE USED OR POSSESSED

SAME

4. NAME OF PERSON TO BE CONTACTED ABOUT THIS APPLICATION Tom Dickinson, Consultant, NMA-
Medical Physics Consultation, 1827 Belt Way Dr., St. Louis MO 63114 TELEPHONE NUMBER (314) 427-5833

SUBMIT ITEMS 5 THROUGH 11 ON 8 1/2 x 11" PAPER. THE TYPE AND SCOPE OF INFORMATION TO BE PROVIDED IS DESCRIBED IN THE LICENSE APPLICATION GUIDE

5. RADIOACTIVE MATERIAL
a. Element and mass number, b. chemical and/or physical form, and c. maximum amount which will be possessed at any one time

6. PURPOSE(S) FOR WHICH LICENSED MATERIAL WILL BE USED

7. INDIVIDUAL(S) RESPONSIBLE FOR RADIATION SAFETY PROGRAM AND THEIR TRAINING AND EXPERIENCE

8. TRAINING FOR INDIVIDUALS WORKING IN OR FREQUENTING RESTRICTED AREAS

9. FACILITIES AND EQUIPMENT.

10. RADIATION SAFETY PROGRAM

11. WASTE MANAGEMENT.

12. LICENSEE FEES (See 10 CFR 170 and Section 170.31)
FEE CATEGORY 7C AMOUNT ENCLOSED \$120

13. CERTIFICATION (Must be completed by applicant) THE APPLICANT UNDERSTANDS THAT ALL STATEMENTS AND REPRESENTATIONS MADE IN THIS APPLICATION ARE BINDING UPON THE APPLICANT. THE APPLICANT AND ANY OFFICIAL EXECUTING THIS CERTIFICATION ON BEHALF OF THE APPLICANT, NAMED IN ITEM 2, CERTIFY THAT THIS APPLICATION IS PREPARED IN CONFORMITY WITH TITLE 10, CODE OF FEDERAL REGULATIONS, PARTS 30, 32, 33, 34, 35, AND 40 AND THAT ALL INFORMATION CONTAINED HEREIN, IS TRUE AND CORRECT TO THE BEST OF THEIR KNOWLEDGE AND BELIEF.
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.

SIGNATURE—CERTIFYING OFFICER

TYPED/PRINTED NAME
Charlotte Lehmann

TITLE
Vice President of Ancillary

DATE
5/30/89

X Charlotte Lehmann

Services

9001300020 890629
REG LIC 30
24-24518-01 PDR

CONTROL NO. 87455

TYPE OF FEE

FILE LOG

FEE CATEGORY

COMMENTS

APPROVED BY

AMOUNT RECEIVED

CHECK NUMBER

RECEIVED

JUN 01 1989

REGION III

DATE

and June 3 7C

\$120 007920

CRP
6/1/89

ITEM 7.1

Additional individuals to be added as Authorized Users for Medical Use.

AUTHORIZED USER

AUTHORIZATION

1) Young Sik Kho, M.D.

35.100, 35.200, 35.300,
35.400

Please refer to the attached Radioactive Material License Nos. 12-03755-01 and IL-00173-01 for evidence of training and experience.

2) Nalini Bidani, M.D.

35.100, 35.200, 35.300

Dr. Bidani is certified in nuclear medicine by the American Board of Nuclear Medicine (1976 Cert. #03922). She was Assistant Professor Radiology, Section of Nuclear Medicine (1979-1986) at the University of Chicago, Chicago, Illinois.

ITEM 7.3

Edward P. Feutz, M.D., is the Radiation Safety Officer at this facility.

Item #7
Page 1 of 1
Prepared: 5/1/89
Lic. #24-24518-01

STATE OF ILLINOIS
DEPARTMENT OF NUCLEAR SAFETY

RADIOACTIVE MATERIAL LICENSE

DIVISION OF NUCLEAR MATERIALS
1035 OUTER PARK DRIVE
SPRINGFIELD, ILLINOIS 62704

Pursuant to the Illinois Radiation Protection Act and the rules and regulations in 22 Illinois Administrative Code promulgated thereunder, 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 and transfer radioactive material(s) listed herein; and to use such radioactive material(s) for the purpose(s) and at the place(s) designated below. This license is subject to all applicable rules, regulations and orders of the Illinois Department of Nuclear Safety now or hereafter in effect and to any conditions specified in the license.

LICENSEE The Memorial Hospital 1501 North Park Drive Belleville, Illinois 62223 Attention: Harry R. Maier, President	LICENSE NUMBER IL-00173-01	EXPIRATION DATE June 30, 1992
	AMENDMENT NUMBER <input type="checkbox"/> INITIAL LICENSE 15	REFERENCE NUMBER 9067414
	<input type="checkbox"/> Initial License <input type="checkbox"/> Amendment through Amendment constitute a complete license. <input checked="" type="checkbox"/> PREVIOUS AMENDMENTS ARE VOID.	

RADIOISOTOPE		MAXIMUM NUMBER OF SOURCES	MAXIMUM ACTIVITY OR QUANTITY PER SOURCE	SEALED SOURCE IDENTIFICATION	DEVICE, SOURCE HOLDER OR CONTAINER	AUTHORIZED USE
ELEMENT	MASS NO.			CHEMICAL and/or PHYSICAL FORM		
Co	57	Total	200 mCi	Prepackaged Kits		In vitro studies
Any in groups I or II		Total as necessary		Any radiopharmaceutical listed in Groups I or II, 32 Ill. Adm. Code 330. App.C		Any diagnostic procedure listed in Groups I or II, 32 Ill. Adm. Code 330. App. C
Co	57	Any	2 mCi	Sealed sources (Nuclear Associates Model 67-294)		Calibration, reference, flood studies or anatomical marking
Co	57	Any	1 mCi	Sealed sources (Picker Model 615205)		Testing and calibration of radiation detection instrumentation.
Co	57	Any	5 mCi	Sealed sources (New England Nuclear Model NES-206)		Quality control testing of dose calibrator

- Radioactive material shall be used only at the licensee's address stated above.
- Radioactive material is authorized for use by, or under the supervision of, the following individual(s) for the materials and uses indicated:

NAME	USES
William Kamm, M.D.	All
Young Sik Kho, M.D.	All
Salvatore Conti, M.D.	All
Mary W. Conti, M.D.	All
Charles L. Mitchell, M.D.	All
Lawrence Taylor, M.D.	In Vitro studies
Edwin Paul Linnette, Ph.D.	In Vitro studies

μCi-Microcurie; mCi-Millicurie; Ci-Curie

BY: <i>Steven C. Collins</i>	DATE June 24, 1987	PAGE 1	OF 4	PAGES
------------------------------	-----------------------	-----------	---------	-------

Steven C. Collins, Chief
L 473-0058

MATERIALS LICENSE

Amendment Number 40

Under the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, of Federal Regulations, Chapter 1, Parts 30, 31, 32, 33, 34, 35, 36, 40 and 70, and in reliance on statements and representations made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s); and to such byproduct and source material. 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 Nuclear Regulatory Commission now or hereafter in effect and to any conditions specified below.

<p>Licensee</p> <p>The Memorial Hospital Department of Nuclear Medicine</p> <p>501 North Park Drive Delewareville, Illinois 62223</p>	<p>In accordance with application dated September 22, 1982,</p> <p>3. License number 12-03755-01 is amended in its entirety to read as follows:</p> <hr/> <p>4. Expiration date January 31, 1988</p> <hr/> <p>5. Docket or Reference No. 030-01452</p>
-----------------------------------------------------------------------------------------------------------------------------------------------	--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

Product, source, and/or special nuclear material	7. Chemical and/or physical form	8. Maximum amount that licensee may possess at any one time under this license
Byproduct material listed in Groups I and II of Schedule A, Section 35.100 of 10 CFR 35	A. Any radiopharmaceutical 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.
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. 4 curies of each byproduct material authorized in Subitem 6.B.
Byproduct material listed in Group IV of Schedule A, Section 35.100 of 10 CFR 35	C. Any radiopharmaceutical listed in Group IV of Schedule A, Section 35.100 of 10 CFR 35	C. As necessary for uses authorized in Subitem 9.C.
Byproduct material listed in Group VI of Schedule A, Section 35.100 of 10 CFR 35	D. Any sealed source listed in Group VI of Schedule A, Section 35.100 of 10 CFR 35	D. 1 curie total for all sources authorized in Subitem 6.D.

MATERIALS LICENSE
SUPPLEMENTARY SHEET

License number

12-03755-01

Docket or Reference number

030-01452

Amendment Number 40

Continued from Page 1

Byproduct, source, and/or special nuclear material	7. Chemical and/or physical form	8. Maximum amount that licensee may possess at any one time
----------------------------------------------------	----------------------------------	-------------------------------------------------------------

Xenon-133

E. Gas or gas in solution that is the subject of an active (i.e., not withdrawn or terminated) "New Drug Application" (NDA) approved by FDA or an active (i.e., not withdrawn, terminated or on "clinical hold") "Notice of Claimed Investigational Exemption for a New Drug" (IND) that has been accepted by FDA

E. 2.5 curies

Any byproduct material listed in Section 31.11(a) of 10 CFR 31

F. Prepackaged kits

F. 3 millicuries of each byproduct material authorized in Subitem 6.F.

Iodine-131

G. Iodide

G. 250 millicuries

Uranium (Depleted in Uranium-235)

H. Cadmium Plated Metal

H. 159 kilograms

Authorized use

Any diagnostic procedure listed in Groups I and II of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.

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.

Any therapeutic procedure listed in Group IV of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.

Any procedure listed in Group VI of Schedule A, Section 35.100 of Title 10, Code of Federal Regulations.

Blood flow studies. Pulmonary function studies.

In vitro studies.

For treatment of thyroid carcinoma.

For use as shielding in a Varian Clinac 6X Medical Linear Accelerator.

CONTROL NO. 87455

MATERIALS LICENSE
SUPPLEMENTARY SHEET

License number	12-03755-01
Docket or Reference number	030-01452
Amendment Number 40	

Continued from Page 2

CONDITIONS

0. Licensed material shall be used only at the licensee's facilities located at 4501 North Park Drive, Belleville, Illinois.
1. 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."
2. Licensed material listed in Item 6 above is authorized for use by, or under the supervision of, the following individual(s) for the materials and uses indicated:

Mary W. Conti, M.D.	All
William Kamm, M.D.	Groups I, II, III, IV and VI Xenon-133 <u>In vitro studies</u>
Young Sik Kho, M.D.	Groups I, II, III, IV and VI Xenon-133 <u>In vitro studies</u>
Salvatore Conti, M.D.	Groups I, II, III and IV Xenon-133 <u>In vitro studies</u>
Charles L. Mitchell, M.D.	Groups I, II and III Xenon-133 Iodine-131 for treatment of hyperthyroidism <u>In vitro studies</u>
Lawrence Taylor, M.D.	Types, quantities and uses of byproduct material listed in 10 CFR 31.11 and 10 CFR 35.31
Edwin Paul Lennette, Ph.D.	<u>In vitro studies</u>
Keith L. Mullenger, M.D.	Groups I, II and III Xenon-133 Iodine-131 for treatment of hyperthyroidism <u>In vitro studies</u>
3. Licensed 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.

CONTROL NO. 87455