NRC FORM 374
(7-94)

U.S. NUCLEAR REGULATORY COMMISSION

MATERIALS LICENSE

Amendment No. 31

Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 36, 39, 40, and 70, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). 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.

302061

Licensee	
1. Grim-Smith Hospital and Clinic, Inc. d/b/a Northeast Regional Medical Center	In accordance with letters dated November 15, 1996 and December 2, 1996 3. License Number 24-05245-01 is amended in its entirety to read as follows:
2. 800 West Jefferson Kirksville, MO 63501	4. Expiration Date November 30, 2000
	5. Docket or Reference No. 030-02332

6. Byproduct, 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
A. Any byproduct material identified in 10 CFR 35.100	A. Any radiopharmaceutical identified in 10 CFR 35.100	A. As needed
B. Any byproduct material identified in 10 CFR 35.200	B. Any radiopharmaceutical identified in 10 CFR 35.200 (excluding xenon-133)	B. As needed
C. Any byproduct material identified in 10 CFR 35.300	C. Any radiopharmaceutical identified in 10 CFR 35.300	C. As needed (not to exceed 1 curie of I-131)
D. Any byproduct material identified in 10 CFR 35.500	D. Sealed sources identified in 10 CFR 35.500	D. As needed
E. Any byproduct material identified in 10 CFR 31.11	E. Prepackaged Kits	E. As needed

0/1

280027

[redacted]

2 mL 30 SD

MATERIALS LICENSE
SUPPLEMENTARY SHEET

License Number
24-05245-01

Docket or Reference Number
030-02332

Amendment No. 31

9. Authorized Use:

A. Medical use described in 10 CFR 35.100.

B. Medical use described in 10 CFR 35.200 (excluding xenon-133).

C. Medical use described in 10 CFR 35.300.

D. Medical use described in 10 CFR 35.500 in devices which have been evaluated and approved for licensing purposes by the U.S. Nuclear Regulatory Commission or an Agreement State.

E. In vitro studies.

CONDITIONS

10. Location of Use: 800 West Jefferson, Kirksville, Missouri.

11. Radiation Safety Officer: Paul M. Williams, D.O.

12. Licensed material listed in Item 6 above is only authorized for use by, or under the supervision of, the following individuals for the materials and uses indicated:

Authorized Users	Material and Use
A. Paul M. Williams, D.O.	10 CFR 35.100, 35.200 (excluding xenon-133), 35.300, 35.500, and 31.11.
B. Michael K. Willman, D.O.	10 CFR 35.100, 35.200 (excluding xenon-133), 35.300, 35.500, and 31.11.
C. G. David Runyon, D.O.	10 CFR 35.100, 35.200 (excluding xenon-133), 35.300, 35.500, and 31.11.

13. The licensee shall collect spent aerosol in a shielded trap and, for reusable traps, monitor the trap effluent with an air contamination monitor that will be checked regularly according to the manufacturer's instruction.

14. The licensee shall not directly vent spent aerosols to the atmosphere and therefore no effluent estimation is necessary.

MATERIALS LICENSE
SUPPLEMENTARY SHEET

License Number 24-05245-01
Docket or Reference Number 030-02332
Amendment No. 31

15. Except as specifically provided otherwise in this license, the licensee shall conduct its program in accordance with the statements, representations, and procedures contained in the documents, including any enclosures, listed below, except for minor changes in the medical use radiation safety procedures as provided in 10 CFR 35.31. The Nuclear Regulatory Commission's regulations shall govern unless the statements, representations, and procedures in the licensee's application and correspondence are more restrictive than the regulations.

A. Application dated April 30, 1990; and

B. Letters dated August 22, 1990 (with attachments), November 15, 1996 and December 2, 1996.

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date December 13, 1996

By Colleen C. Casey
Nuclear Materials Licensing Branch, Region III

BETWEEN:

License Fee Management Branch, ARM
and
Regional Licensing Sections

(FOR LFMS USE)
INFORMATION FROM LTS

Program Code: 02120
Status Code: 0
Fee Category: 7C
Exp. Date: 20001130
Fee Comments:
Decom Fin Assur Reqd: N

S6

LICENSE FEE TRANSMITTAL

A. REGION

1. APPLICATION ATTACHED
   Applicant/Licensee: GRIM-SMITH HOSPITAL & CLINIC, INC.
   Received Date: 961119
   Docket No: 3002332
   Control No.: 302061
   License No.: 24-05245-01
   Action Type: Amendment

2. FEE ATTACHED
   Amount: 440
   Check No.: 38266

3. COMMENTS

Signed D. Hersey
Date 11-21-96

96 NOV 25 AM 11:25

B. LICENSE FEE MANAGEMENT BRANCH (Check when milestone 03 is entered (✓))

1. Fee Category and Amount: 7C $440

2. Correct Fee Paid. Application may be processed for:
   Amendment ✓
   Renewal
   License

3. OTHER

Signed SC
Date 11/26/96

NOV 29 1996

Log NOV II III
Remitter
Check No. 38266
Amount $440
Fee Category 7C
Type of Fee AMD
Date Check Rec'd 11/25/96
Date Completed 11/26/96
By: SC

LAW OFFICES

SHOOK, HARDY & BACON L.L.P.

40 CORPORATE WOODS, 6TH FLOOR
9401 INDIAN CREEK PARKWAY
P.O. BOX 25128
OVERLAND PARK, KANSAS 66225-5128
TELEPHONE (913) 451-6060 ▪ FACSIMILE (913) 451-6879

A LIMITED LIABILITY
PARTNERSHIP INCLUDING
PROFESSIONAL CORPORATIONS

KANSAS CITY, MISSOURI
HOUSTON, TEXAS
LONDON, ENGLAND
ZURICH, SWITZERLAND
MILAN, ITALY

November 15, 1996

VIA FEDERAL EXPRESS

U.S. Nuclear Regulatory Commission
Chief, Materials Licensing Branch
801 Warrenville Road
Lisle, Illinois 60532-4351

Re: **Request for Transfer of Materials License #24-05245-01**

Dear Sirs:

Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center (Grim-Smith) and Kirksville Osteopathic Medical Center (KOMC) propose to merge and consolidate operations on or about December 2, 1996. After the closing of the transaction, the operating assets of both hospitals will be owned and operated by a single corporate entity, Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center, which will then operate two facilities under a consolidated license: (1) The Patterson Campus, 112 East Patterson, Kirksville, Missouri 63501; and, (2) The Jefferson Campus, 800 West Jefferson, Kirksville, Missouri 63501.

Both Grim-Smith and KOMC currently hold Materials Licenses with the Nuclear Reglatory Commission. Pursuant to Attachment 1 IN89-25, Rev.1, December 7, 1994, enclosed please find our client's request for transfer of the KOMC Materials License #24-05245-01 to Grim-Smith. We realize that this request is not being made 90 days before the proposed transfer (as specified by regulation); however, due to the nature of the merger, we were unable to complete the application until now. We would sincerely appreciate your agency's assistance in reviewing this request for transfer as soon as possible in order to facilitate continuity of patient care. We are enclosing copies of the Materials Licenses of both Grim-Smith and KOMC for your convenience.

Thank you for your assistance with this request. Please do not hesistate to contact me should you have any questions or comments.

Sincerely,

Carol A. Poindexter

CAPds
Enclosures

0072804.01

Pm: 11-18-96

GRIM-SMITH

H O S P I T A L

112 E. Patterson, Kirksville, MO 63501
(816) 665-7241
FAX: (816) 665-0302

November 11, 1996

US Nuclear Regulatory Commission
Region III
801 Warrenville Road
Lisle, Illinois 60532-4351

Re: License # 24-05245-01

Dear Sirs:

Attached is an application for Change of Ownership of the above referenced By-Product Materials (Medical Use) License issued to Kirksville Osteopathic Medical Center, 800 West Jefferson, Kirksville, MO 63501.

We anticipate that the change of ownership of the facility will take place on November 26, 1996. Due to the short time frame under which we are operating we would greatly appreciate any help you could give us in getting the change of license ownership authorized as soon as possible.

Also attached is check for $440 issued to the US Nuclear Regulatory Commission to cover the cost of this change.

If there are any questions regarding this application, please contact: Mark Collins, Departmental Administrator, Grim-Smith Hospital (816) 785-3714, or our consulting physicist, Jon J. Erickson (816) 390-9011, or our legal counsel, Carol Poindexter (913) 451-6060.

Thank you for your help.

Sincerely,

Charles M. Boughton
Vice President-Finance

CMB/ds

Application For Change of Ownership

The following is an application of authorization to change ownership of By-Product License #24-05245-01 currently held by Kirksville Osteopathic Medical Center, 800 West Jefferson, Kirksville, MO 63501.

The item number refers to the corresponding items listed in Attachment I of IN 89-25, Rev 1, December 7, 1994.

The current owner of this license is the Kirksville Osteopathic Medical Center (KOMC). The new owner is to be:

Grim-Smith Hospital & Clinic, Inc.
d/b/a Northeast Regional Medical Center
112 E. Patterson Avenue, Kirksville, MO 63501

Item 1. Current name of licensed organization:
Kirksville Osteopathic Medical Center
Future name of licensed organization:
Grim-Smith Hospital & Clinic, Inc.
d/b/a Northeast Regional Medical Center

Item 2. New license contact:
Mark Collins
(816) 785-3714

Item 3. Changes in personnel:

Following change of ownership the current operations personnel (CEO, CFO, President, etc.) of Grim-Smith Hospital will have control over, and be responsible for the daily operation of the Northeast Regional Medical Center.

There will be no change in the personnel responsible for the radiation safety program and use of by-product material. The personnel currently employed by Kirksville Osteopathic Medical Center as Radiation Safety Officer and Authorized Users will remain in those positions following the change of ownership.

Item 4. Kirksville Osteopathic Medical Center will not remain in non-licensed business without the license.

Item 5. Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center (Grim-Smith) and Kirksville Osteopathic Medical Center (KOMC) propose to consummate a merger/consolidation transaction whereby the operations of

those entities will be consolidated into Grim-Smith on or about November 26, 1996. After the Closing of the transaction, the operating assets of both hospitals will be owned and operated by a single corporate entity, Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center, which will then operate both facilities under a consolidated license: (1) The Patterson Campus, 112 East Patterson, Kirksville, Missouri 63501; and, (2) The Jefferson Campus, 800 West Jefferson, Kirksville, Missouri 63501.

Item 6. There will be no changes in the organization (other than upper management), location, equipment, or procedures following the change in ownership.

Item 7. There will be no change in the use, possession, location or storage of the licensed material.

Item 8. There are no changes in organization, location, facilities, equipment, procedures, or personnel that would require a license amendment without the change in ownership.

Item 9. All surveillance items and records will be current at the time of transfer. All of the surveillance requirements and records are current at the present time.

Item 10. Since licensed activity will continue at the same location, all records concerning safe and effective decommissioning of the facility pursuant to 10CFR 30.35(g), 40.36(f), 70.25(g), and 72.30(d); public dose, waste disposal by release to sewers, incineration, radioactive spills, and on-site burials have been transferred to the new licensee.

Item 11. The facility is currently operating within the guidelines of good radiation safety practices and there is no radioactive contamination present.

Item 12. Not applicable.

Item 13. Covered in attached Letter of Confirmation from Grim-Smith Hospital.

Item 14. Covered by the attached Letters of Agreement from Grim-Smith Hospital and Kirksville Osteopathic Medical Center

Submitted by:

Charles M. Boughton,
Vice President-Finance

0072271.01

GRIM-SMITH

H O S P I T A L

112 E. Patterson, Kirksville, MO 63501
(816) 665-7241
FAX: (816) 665-0302

November 11, 1996

TO WHOM IT MAY CONCERN:

Re: Item 13: Attachment 1, IN89-25, Rev. 1, December 7, 1994

By this letter we confirm that we agree to abide by all commitments and representations previously made to NRC by Kirksville Osteopathic Medical Center. These include but are not limited to: maintaining decommissioning records required by 10CFR30.35(g); implementing decontamination activities and decommissioning of the site; and completing corrective actions for open inspection items and enforcement actions.

We further confirm that we will abide by all provisions under Item 13: Attachment 1, IN89-25, Rev. 1, December 7, 1994. In regard to financial assurance arrangements to fund decommissioning, it is our understanding that the level of operation of this facility does not require a formal funding program for decommissioning.

With regard to open inspection items, we are in possession of a written commitment (attached) from Kirksville Osteopathic Medical Center indicating that there are no open inspection items.

Sincerely,

Charles M. Boughton
Vice President-Finance

CMB/ds

Kirksville Osteopathic
Medical Center

Northeast Missouri's premier
health care provider since 1905!

November 15, 1996

TO WHOM IT MAY CONCERN:

RE: Kirksville Osteopathic Medical Center Nuclear Regulatory Commission ("NRC")
Materials License No. 24-05245-01; NRC Information Notice 39-25, Rev. 1,
Attachment 1, Item 14, December 7, 1994

We hereby confirm that Kirksville Osteopathic Medical Center ("KOMC"), which holds the referenced NRC Materials License, is a party to a merger/consolidation transaction whereby the operations and assets of KOMC will be acquired by Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center.

In conjunction with that transaction, and pursuant to Item 14 of Attachment 1 of IN89-25, Rev. 1, KOMC agrees to the change in ownership and control to Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center of: 1) the material regulated under the Materials License; 2) the activities authorized by the Materials License; and 3) the conditions of transfer as described in Attachment 1 to IN89-25, Rev.1.

Grim-Smith Hospital & Clinic, Inc. has been notified that there are no open Materials License deficiencies documented by NRC inspections and that Grim-Smith Hospital & Clinic, Inc., is responsible for possible enforcement actions resulting from any deficiencies.

Sincerely

Zell Cook
Chief Executive Officer

MATERIALS LICENSE

Amendment No. 30

Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 39, 40 and 70, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). 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.

Licensee	
1. Kirksville Osteopathic Medical Center 2. 800 West Jefferson Kirksville, MO 63501	In accordance with application dated April 30, 1990. 3. License number 24-05245-01 is renewed in its entirety to read as follows: 4. Expiration date November 30, 1995 5. Docket or Reference No. 030-02332

6. Byproduct, 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
A. Any byproduct material identified in 10 CFR 35.100	A. Any radiopharmaceutical identified in 10 CFR 35.100	A. As needed
Any byproduct material identified in 10 CFR 35.200	B. Any radiopharmaceutical identified in 10 CFR 35.200 (excluding Xenon-133)	B. As needed
C. Any byproduct material identified in 10 CFR 35.300	C. Any radiopharmaceutical identified in 10 CFR 35.300	C. As needed
D. Any byproduct material identified in 10 CFR 35.500	D. Sealed sources identified in 10 CFR 35.500	D. As needed
E. Any byproduct material identified in 10 CFR 31.11	E. Prepackaged Kits	E. As needed

9. Authorized Use:

A. Medical use described in 10 CFR 35.100.

B. Medical use described in 10 CFR 35.200 (excluding Xenon-133).

C. Medical use described in 10 CFR 35.300.

D. Medical use described in 10 CFR 35.500 in devices which have been evaluated and approved for licensing purposes by the U.S. Nuclear Regulatory Commission or an Agreement State.

E. In vitro studies.

CONDITIONS

10. Location of Use: 800 West Jefferson, Kirksville, Missouri.

11. Radiation Safety Officer: Paul M. Williams, D.O.

12. Authorized Users:

A. Paul M. Williams, D.O., for material in 10 CFR 35.100, 35.200, (excluding Xenon-133), 35.300, 35.500, and 31.11.

B. Michael K. Willman, D.O., for material in 10 CFR 35.100, 35.200, (excluding Xenon-133), 35.300, 35.500, and 31.11.

C. G. David Runyon, D.O., for material in 10 CFR 35.100, 35.200, (excluding Xenon-133), 35.300, 35.500, 31.11.

13. Pursuant to Title 10, Chapter 1, Code of Federal Regulations, Part 40, "Domestic Licensing of Source Material," the licensee is authorized to possess, use, transfer, and import up to 999 kilograms of depleted uranium contained as shielding material in the molybdenum-99/technetium-99m generators authorized by this license.

14. The licensee shall collect spent aerosol in a shielded trap and, for reusable traps, monitor the trap effluent with an air contamination monitor that will be checked regularly according to the manufacturer's instruction.

15. The licensee shall not directly vent spent aerosols to the atmosphere and therefore no effluent estimation is necessary.

16. The licensee shall maintain records of information important to safe and effective decommissioning at the address in Condition 10. per the provisions of 10 CFR 30.35(g) until this license is terminated by the Commission.

MATERIALS LICENSE
SUPPLEMENTARY SHEET

License number
24-05245-01

Docket or Reference number
030-02332

Amendment No. 30

17. This license is based on the licensee's statements and representations listed below:

A. Application dated April 30, 1990; and

B. Letter dated August 22, 1990 (with attachments)

For the U.S. Nuclear Regulatory Commission

Date: September 6, 1990

By Robert D. Gattozzi
Materials Licensing Section, Region 1

NRC FORM 374 (7-84)

U.S. NUCLEAR REGULATORY COMMISSION

MATERIALS LICENSE

Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 36, 39, 40, and 70, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). 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.

Licensee

1. Grim-Smith Hospital

2. 112 East Patterson
Kirksville, MO 63501

3. License Number 24-26702-01

4. Expiration Date April 30, 2001

5. Docket or Reference No. 030-34073

6. Byproduct, 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
A. Any byproduct material identified in 10 CFR 35.100	A. Any radiopharmaceutical identified in 10 CFR 35.100	A. As needed
B. Any byproduct material identified in 10 CFR 35.200	B. Any radiopharmaceutical identified in 10 CFR 35.200 (excluding xenon-133)	B. As needed
C. Any byproduct material identified in 10 CFR 35.300	C. Any radiopharmaceutical identified in 10 CFR 35.300	C. 1 Curie
D. Any byproduct material identified in 10 CFR 31.11	D. Prepackaged Kits	D. As needed

COPY

NRC FORM 374A (7-84) U.S. NUCLEAR REGULATORY COMMISSION

License Number 24-26702-01

Docket or Reference Number 030-34073

MATERIALS LICENSE
SUPPLEMENTARY SHEET

9. Authorized Use:

A. Medical use described in 10 CFR 35.100.

B. Medical use described in 10 CFR 35.200 (excluding xenon-133).

C. Medical use described in 10 CFR 35.300.

D. In vitro studies.

CONDITIONS

10. Licensed material shall be used only at the licensee's facilities located at 112 East Patterson, Kirksville, Missouri.

11. Radiation Safety Officer: Paul M. Williams, D.O.

12. Licensed material listed in Item 6 above is only authorized for use by, or under the supervision of, the following individuals for the materials and uses indicated:

Authorized Users	Material and Use
A. Paul M. Williams, D.O.	10 CFR 35.100, 35.200 (excluding xenon-133), 35.300 and 31.11.
B. Michael K. Willman, D.O.	10 CFR 35.100, 35.200 (excluding xenon-133), 35.300 and 31.11.

13. The licensee may not possess and use materials authorized in Items 6, 7, and 8 until:

A. The licensee has constructed the facilities and obtained the equipment described in the application and supporting documentation; and

B. The U. S. Nuclear Regulatory Commission, Region III, ATTN: Chief, Materials Licensing Branch, 801 Warrenville Road, Lisle, IL 60532-4351 has been notified that activities authorized by the license will be initiated.

COPY

NRC FORM 374A (7-94) U.S. NUCLEAR REGULATORY COMMISSION

MATERIALS LICENSE
SUPPLEMENTARY SHEET

License Number: 24-26702-01

Docket or Reference Number: 030-34073

14. Within 30 days of the date of a decision not to complete the facility, acquire equipment, or possess and use authorized material, the licensee must notify the Commission in writing, of the decision.

15. Except as specifically provided otherwise in this license, the licensee shall conduct its program in accordance with the statements, representations, and procedures contained in the documents, including any enclosures, listed below, except for minor changes in the medical use radiation safety procedures as provided in 10 CFR 35.31. The Nuclear Regulatory Commission's regulations shall govern unless the statements, representations, and procedures in the licensee's application and correspondence are more restrictive than the regulations.

A. Application dated February 15, 1996 (except Quality Management Program); and

B. Letters dated March 15, 1996 (except Quality Management Program) and March 28, 1996.

FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date 9 April 1996

By William L. Kinkead
Materials Licensing Branch, Region III

COPY

DEC 16 1996

Chuck Boughton, Vice President
Grim-Smith Hospital and Clinic, Inc.
d/b/a Northeast Regional Medical Center
800 West Jefferson
Kirksville, MO 63501

Dear Mr. Boughton:

Enclosed is Amendment No. 31 to your NRC Material License No. 24-05245-01 in accordance with your request.

Please review the enclosed document carefully and be sure that you understand all conditions. If there are any errors or questions, please notify the U.S. Nuclear Regulatory Commission, Region III office at (630) 829-9887 so that we can provide appropriate corrections and answers.

A. In preparing this amendment, we took the opportunity to update and reformat your license as discussed below. These changes were essentially administrative modifications only that should have no effect on your licensed program.

Subitems A.1., A.2., A.3. and A.4. of this letter were discussed in telephone conversations between Colleen Casey and Mark Collins on November 27, 1996.

1. Please note that, at this time, we changed the expiration date in item no. 4 of your license to reflect the one-time extension of your license, in accordance with 10 CFR 30.36(a)(2), copy enclosed. You should have received additional correspondence from us concerning this regulation and its effects on your license.

2. We deleted Condition No. 13., as it appeared on Amendment No. 30, which authorized the possession, use, transfer and import of up to 999 kilograms of depleted uranium (DU) contained as shielding material for generators.

   We have been advised that you may be subject to additional annual fees if you retained the DU authorization on your license, although it is highly unlikely that your licensed program would need this authorization. In light of these considerations, Mr. Collins directed us to remove the DU authorization from your license in a letter dated December 2, 1996.

3. We deleted Condition No. 16., as it appeared on Amendment No. 30, because the regulations in 10 CFR 30.35(g) contain the same provision. Therefore, this Condition is no longer necessary.

302061

4. At this time, we limited your authorization for iodine-131, as listed in 10 CFR 35.300 and Item 8.C., in order to preclude your having to file an emergency plan, per 10 CFR 30.32(i) and 30.72, enclosed. Your total possession limit for iodine-131 is one curie total and includes waste activity also. Mr. Collins advised us of the possession limit you wish to have for iodine-131 in the letter dated December 2, 1996.

5. We reformatted Condition Nos. 12. and 17. to conform with our current licensing style.

6. If your understanding differs from ours in any of these matters, or if you have questions concerning this amendment, please contact Colleen C. Casey at (630) 829-9841.

B. This also refers to your letters dated November 15, 1996 (including two attached letters dated November 11, 1996) and December 2, 1996, concerning your institution's change in ownership.

Based upon statements submitted in your letters regarding this change, NRC has no objection to the acquisition of Kirksville Osteopathic Medical Center by Grim-Smith Hospital & Clinic, Inc., d/b/a Northeast Regional Medical Center. We have amended your license to reflect the change in your medical center's name.

However, please be reminded that 10 CFR 30.34(b), enclosed, states, "No license issued or granted pursuant to the regulations in this part, Parts 31 through 36, and 39, nor any right under a license shall be transferred, assigned or in any manner disposed of, either voluntarily or involuntarily, directly or indirectly, through transfer of control of any license to any person, unless the Commission shall, after securing full information, find that the transfer is in accordance with the provisions of the Act and shall give its consent in writing."

Although it is not NRC's intent to interfere with the business decisions of licensees, it is necessary for licensees to provide timely notification to NRC whenever such decisions could involve changes in the corporate structure responsible for management oversight, control, or radiological safety of licensed materials. Full information on changes of ownership or control of licensed activities should be submitted to the appropriate NRC regional or Headquarters office, 90 days prior to the proposed action.

The purpose of such notification is to allow NRC to assure that: (1) radioactive materials are possessed, used, owned, or controlled only by persons who have valid NRC licenses; (2) materials are properly handled and secured; (3) persons using such materials are capable, competent, and committed to implement appropriate radiological controls; (4) licensees provide adequate financial assurance for compliance with NRC requirements; and (5) public health and safety are not

compromised by the use of such materials. Although the burden of notification is on the existing licensee, it may still be necessary for the transferee to provide supporting information or to independently coordinate the change in ownership or control with the appropriate NRC office.

NRC licensees planning to transfer ownership, to change the corporate status, or to change control of licensed activities are required to provide sufficient prior notice and full information about the change to NRC, and to obtain written consent from the Commission before the transfer. Failure to comply with this requirement may adversely affect the public health and safety and interfere with NRC's ability to inspect licensed activities. Cases where change of ownership or control has occurred without prior written consent from NRC will be treated as noncompliance with the provisions of 10 CFR 30.34 and will be referred to the inspection staff and/or Office of Investigations, as appropriate.

The failure to receive required NRC approval prior to a change in ownership or control of licensed activities can be considered a violation of NRC requirements and may warrant escalated enforcement action, to include civil penalties and orders, Willful failure to obtain prior NRC approval of the transfer may result in referrals to the Department of Justice for consideration of criminal prosecution. Further information and guidance in these matters is contained in the enclosed Information Notice 89-25, Rev. 1, dated December 7, 1994.

C. Please be advised that your license expires at the end of the day, in the month, and year stated in the license. Unless your license has been terminated, you must conduct your program involving byproduct materials in accordance with the conditions of your NRC license, representations made in your license application, and NRC regulations. In particular, note that you must:

1. Operate in accordance with NRC regulations 10 CFR Part 19, "Notices, Instructions and Reports to Workers; Inspections," 10 CFR Part 20, "Standards for Protection Against Radiation," and other applicable regulations.

2. Notify NRC, in writing, within 30 days:

   a. When an authorized user or Radiation Safety Officer permanently discontinues performance of duties under the license or has a name change; or

   b. When the licensee's mailing address changes (no fee is required if the location of byproduct material remains the same).

3. In accordance with 10 CFR 30.36(b) and/or license condition, notify NRC, promptly, in writing, and request termination of the license when you decide to terminate all activities involving materials authorized under the license.

4. Request and obtain a license amendment before you:

   a. Receive or use byproduct material for a clinical procedure permitted under Part 35 but not permitted by your license issued pursuant to this Part;

   b. Permit anyone, except individuals described in 10 CFR 35.13(b), to work as an authorized user under the license;

   c. Change Radiation Safety Officers;

   d. Order byproduct material in excess of the amount, or radionuclide, or form different than authorized on the license;

   e. Add or change the areas of use or address or addresses of use identified in the license application or on the license; or

   f. Change ownership of your organization.

5. Submit a complete renewal application with proper fee or termination request at least 30 days before the expiration date of your license. You will receive a reminder notice approximately 90 days before the expiration date. Possession of byproduct material after your license expires is a violation of NRC regulations. A license will not normally be renewed, except on a case-by-case basis, in instances where licensed material has never been possessed or used.

In addition, please note that NRC Form 313 requires the applicant, by his/her signature, to verify that the applicant understands that all statements contained in the application are true and correct to the best of the applicant's knowledge. The signatory for the application should be the licensee or certifying official rather than a consultant.

You will be periodically inspected by NRC. Failure to conduct your program in accordance with NRC regulations, license conditions, and representations made in your license application and supplemental correspondence with NRC will result in enforcement action against you. This could include issuance of a notice of violation, or imposition of a civil penalty, or an order suspending, modifying or revoking your license as specified in the General Policy and Procedures for NRC Enforcement Actions. Since serious consequences to employees and the public can result from failure to comply with NRC requirements,

prompt and vigorous enforcement action will be taken when dealing with licensees who do not achieve the necessary meticulous attention to detail and the high standard of compliance which NRC expects of its licensees.

Sincerely,

Original Signed By
Colleen C. Casey
Nuclear Materials Licensing Branch

License No.: 24-05245-01
Docket No.: 030-02332

Enclosures:
1. Amendment No. 31
2. 10 CFR Part 30
3. NRC Form 313
4. IN 89-25, Rev. 1

DOCUMENT NAME: M:\03002332.CL6

To receive a copy of this document, indicate in the box: "C" = Copy without attachment/enclosure "E" = Copy with attachment/enclosure "N" = No copy

OFFICE	DNMS/RIII CCC	C							
NAME	CCASEY:jaw								
DATE	12/3/96								

OFFICIAL RECORD COPY

GRIM-SMITH
HOSPITAL

112 E. Patterson, Kirksville, MO 63501
(816) 665-7241
FAX: (816) 665-0302

December 2, 1996

This document is in response to a telephone conversation and follow-up documents done by Colleen Casey on November 27, 1996.

I. Grim-Smith Hospital & Clinic, Inc. does not wish to terminate its license at 112 East Patterson, license #24-26702-01. There has been no change of ownership and there has been no name change.

II.

A. 1. We as Grim-Smith Hospital & Clinic, Inc. d/b/a Northeast Regional Medical Center commit to abide by all constraints, conditions, requirements, representations, and commitments identified in the existing license #24-05245-0. Please refer to Chuck Boughton's letter dated November 11, 1996 regarding item 13.

2. The new contact person is Mark S. Collins. His position by title is Administrative Director of Clinical Services. He can be reached at 816-785-3714.

3. Grim-Smith Hospital & Clinic, Inc. d/b/a Northeast Regional Medical Center confirm as the new owners to accept full liability for the site.

B. Grim-Smith Hospital & Clinic, Inc. d/b/a Northeast Regional Medical Center is requesting that condition #13 be removed from the license.

C. The possession limit Grim-Smith Hospital & Clinic, Inc. d/b/a Northeast Regional Medical Center wishes to have for iodine-131 is one Curie.

D. Grim-Smith Hospital & Clinic, Inc. d/b/a Northeast Regional Medical Center do not wish to change the mailing address on the license. The correct address is:

800 West Jefferson.
Kirksville MO 63501

Sincerely

Chuck Boughton
Vice President

12-3-96

GRIM-SMITH

HOSPITAL

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Kirksville MO 63501

PM; 12-2-96

TELEPHONE CONVERSATION RECORD

BETWEEN

Colleen C. Casey, NMLB Reviewer and Mark Collins of Northeast Regional Medical Center on November 27, 1996 at P.M. (time). Ms. Casey represents the United States Nuclear Regulatory Commission, Region III, Nuclear Materials Licensing Branch, 801 Warrenville Road, Lisle, Illinois 60532-4351.

Control No.: 302061 and 302063 License No.: 24-05245-01 and 24-26702-01

I. To terminate the Grim-Smith license, we will need the following:

A. A completed Form NRC 314;

B. A completed close-out survey, as follows:

Please be advised that we cannot authorize you to release your old nuclear (or facility) medicine space for unrestricted use (even by other members of your staff) until we have received and reviewed a copy of the results of your close-out survey. The survey should consist of exposure rate measurements to show that all sources of radioactive material have been removed, and contamination checks of areas where radioactive materials were used or stored. Average radiation levels associated with surface contamination and removable contamination should not exceed those specified in the enclosed decontamination guide. Please submit the following information with your close-out survey:

a. A diagram of your old facility with survey and wipe test results keyed to specific locations.

b. The name of the person performing the survey.

c. The date the survey was performed.

d. The instrument(s) used for exposure rate measurements and for analysis of the wipes.

e. Background readings and each instruments' efficiency or correction factor.

f. The date that the survey instrument was last calibrated.

*being sent

CONTD NEXT PAGE

OPTIONAL FORM 99 (7-90)

FAX TRANSMITTAL # of pages ▸ 4

To MARK COLLINS	From COLLEEN C. CASEY
Dept./Agency GRIM-SMITH HOSPITAL	Phone # 630-829-9241
Fax # 816-785-3929	Fax # 630-575-1073/1259

NSN 7540-01-317-7368 5099-101 GENERAL SERVICES ADMINISTRATION

C. When a licensee terminates licensed activities or licensed activities are transferred to another licensee, we require that certain records be transferred to the NRC or to a successor licensee to ensure their long-term availability. Please transfer the following records, as were applicable to your program, that pertain to:

1. the decommissioning of your facility (see I.B. above);

2. radiation doses to the public (not dosimetry records for occupationally exposed persons); and

3. waste disposal by release to sewers, incineration, radioactive material spills, and on-site burials. Records documenting the occurrences of spills and the subsequent decontamination activities should be included.

These records should be transferred to the USNRC Region III, 801 Warrenville Road, Lisle, Illinois 60532-4351. No transfers, changes of ownership or license terminations will be authorized unless all records considered important to the safe and effective decommissioning of the facility contained in 10 CFR 30.35(g) and all records concerning public dose and waste disposal have been transferred to the NRC (for license terminations).

II. Change of Ownership for KOMC

To complete our review of your letter dated November 15, 1996, concerning the change of ownership for KOMC, we will need the following additional information:

A. 1. A commitment by the transferee to abide by all constraints, conditions, requirements, representations and commitments identified in the existing license. If not, the transferee must provide a description of its program to ensure compliance with the license and regulations.

2. Please specify the position held by your new contact person, Mark Collins.

3. Please confirm that the new owner accepts full liability for the site.

B. We noted that your license currently contains Condition No. 13., which authorizes the possession, use, transfer and import of up to 999 kilograms of depleted uranium (DU) contained as shielding material in the molybdenum-99/technetium-99m generators authorized by 10 CFR 35.200 on your license.

We have been advised that you may be subject to additional annual fees as a result of the DU authorization on your license. We have also been informed that you may not require the DU authorization, even if you do receive generators. It may be to your benefit to

review the necessity of retaining this authorization with your generator supplier, Radiation Safety Officer, Radiation Safety Committee and senior management.

If you wish, you may direct us to remove the DU authorization from your license in your response to the above items in this letter and we will amend your license accordingly, without an additional fee. Please follow the procedure described below.

C. At this time, we must limit your authorization for iodine-131, as listed in 10 CFR 35.300, in order to preclude your having to file an emergency plan, per 10 CFR 30.32(i) and 30.72, enclosed. Your total possession limit for iodine-131 must be less than 10 curies and will include waste activity also. Please advise us of the possession limit you wish to have for iodine-131.

D. It appears that you wish to change the mailing address for this license to "112 East Patterson Street, Kirksville, MO 63501," which is different than the location where licensed activities will continue to take place. Please confirm if this is so or advise us accordingly if our understanding is incorrect.

15 DAYS RESPONSE- SEND TO MS. CASEY'S ATTENTION AT THE NRC OFFICE ADDRESS GIVEN ABOVE AND REFERENCE CONTROL NOs. 302061/302063.

If you have any questions or require clarification on any of the information stated above, you may contact Colleen C. Casey at (630) 829-9841.

Reviewer's signature: Colleen C. Casey Date: 11/26/96

NRC FORM 314 (6-95) U.S. NUCLEAR REGULATORY COMMISSION

10 CFR 30.36(c)(1)(iv)
10 CFR 40.42(c)(1)(iv)
10 CFR 70.38(c)(1)(iv)

APPROVED BY OMB: NO. 3150-0028 EXPIRES: 06/30/98

ESTIMATED BURDEN PER RESPONSE TO COMPLY WITH THIS MANDATORY INFORMATION COLLECTION REQUEST: 30 MINUTES. THIS SUBMITTAL IS USED BY NRC AS PART OF THE BASIS FOR ITS DETERMINATION THAT THE FACILITY HAS BEEN CLEARED OF RADIOACTIVE MATERIAL BEFORE THE FACILITY IS RELEASED FOR UNRESTRICTED USE. FORWARD COMMENTS REGARDING BURDEN ESTIMATE TO THE INFORMATION AND RECORDS MANAGEMENT BRANCH (T-6 F33), U.S. NUCLEAR REGULATORY COMMISSION, WASHINGTON, DC 20555-0001, AND TO THE PAPERWORK REDUCTION PROJECT (3150-0028), OFFICE OF MANAGEMENT AND BUDGET, WASHINGTON, DC 20503. AN AGENCY MAY NOT CONDUCT OR SPONSOR, AND A PERSON IS NOT REQUIRED TO RESPOND TO, A COLLECTION OF INFORMATION UNLESS IT DISPLAYS A CURRENTLY VALID OMB CONTROL NUMBER.

CERTIFICATE OF DISPOSITION OF MATERIALS

INSTRUCTIONS: ALL ITEMS MUST BE COMPLETED -- PRINT OR TYPE
SEND THE COMPLETED CERTIFICATE TO THE NRC OFFICE SPECIFIED ON THE REVERSE

LICENSEE NAME AND ADDRESS	LICENSE NUMBER
	LICENSE EXPIRATION DATE

A. MATERIALS DATA *(Check one and complete as necessary)*

THE LICENSEE OR ANY INDIVIDUAL EXECUTING THIS CERTIFICATE ON BEHALF OF THE LICENSEE CERTIFIES THAT:
(Check and/or complete the appropriate item(s) below.)

☐ 1. NO MATERIALS HAVE EVER BEEN PROCURED OR POSSESSED BY THE LICENSEE UNDER THIS LICENSE.

OR

☐ 2. ALL ACTIVITIES AUTHORIZED BY THE LICENSE HAVE CEASED AND ALL MATERIALS PROCURED AND/OR POSSESSED BY THE LICENSE NUMBER CITED ABOVE HAVE BEEN DISPOSED OF IN THE FOLLOWING MANNER. *(If additional space is needed, use the reverse side or provide attachments.)*

Describe specific material transfer actions and, if there were radioactive wastes generated in terminating this license, the disposal actions including the disposition of low-level radioactive waste, mixed waste, Greater-than-Class-C waste, and sealed sources, if applicable.

For transfers, specify the date of the transfer, the name of the license recipient, and the recipient's NRC license number or Agreement State name and license number.

If materials were disposed of directly by the licensee rather than transferred to another licensee, licensed disposal site or waste contractor, describe the specific disposal procedures *(e.g., decay in storage)*

B. OTHER DATA

☐ 1. OUR LICENSE HAS NOT YET EXPIRED; PLEASE TERMINATE IT.

2. A RADIATION SURVEY WAS CONDUCTED BY THE LICENSEE TO CONFIRM THE ABSENCE OF LICENSED RADIOACTIVE MATERIALS AND TO DETERMINE WHETHER ANY CONTAMINATION REMAINS ON THE PREMISES COVERED BY THE LICENSE. *(Check one)*

☐ NO *(Attach explanation)*
☐ YES, THE RESULTS *(Check one)*
  ☐ ARE ATTACHED, *or*
  ☐ WERE FORWARDED TO NRC ON *(Date)*

3. THE PERSON TO BE CONTACTED REGARDING THE INFORMATION PROVIDED ON THIS FORM	NAME	TELEPHONE NUMBER *(Include Area Code)*

4. MAIL ALL FUTURE CORRESPONDENCE REGARDING THIS LICENSE TO

CERTIFYING OFFICIAL

I CERTIFY UNDER PENALTY OF PERJURY THAT THE FOREGOING IS TRUE AND CORRECT

PRINTED NAME AND TITLE	SIGNATURE	DATE

WARNING: FALSE STATEMENTS IN THIS CERTIFICATE MAY BE SUBJECT TO CIVIL AND/OR CRIMINAL PENALTIES. NRC REGULATIONS REQUIRE THAT SUBMISSIONS TO THE NRC BE COMPLETE AND ACCURATE IN ALL MATERIAL RESPECTS. 18 U.S.C. SECTION 1001 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 JURISDICTIONS.

FILE CERTIFICATES AS FOLLOWS:

IF YOU ARE A DISTRIBUTOR OF EXEMPT PRODUCTS, SEND TO:

DIVISION OF INDUSTRIAL AND MEDICAL NUCLEAR SAFETY
OFFICE OF NUCLEAR MATERIAL SAFETY AND SAFEGUARDS
U.S. NUCLEAR REGULATORY COMMISSION
WASHINGTON, DC 20555-0001

ALL OTHERS, 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:

LICENSING ASSISTANCE SECTION
NUCLEAR MATERIALS SAFETY BRANCH
U.S. NUCLEAR REGULATORY COMMISSION, REGION I
475 ALLENDALE ROAD
KING OF PRUSSIA, PA 19406-1415

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

NUCLEAR MATERIALS SAFETY SECTION
U.S. NUCLEAR REGULATORY COMMISSION, REGION II
101 MARIETTA STREET NW, SUITE 2900
ATLANTA, GA 30323-0199

IF YOU ARE LOCATED IN:

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

MATERIALS LICENSING SECTION
U.S. NUCLEAR REGULATORY COMMISSION, RECION III
801 WARRENVILLE ROAD
LISLE, IL 60532-4351

ALASKA, ARIZONA, ARKANSAS, CALIFORNIA, COLORADO, HAWAII, IDAHO, KANSAS, LOUISIANA, MONTANA, NEBRASKA, NEVADA, NEW MEXICO, NORTH DAKOTA, OKLAHOMA, OREGON, PACIFIC TRUST TERRITORIES, SOUTH DAKOTA, TEXAS, UTAH, WASHINGTON, OR WYOMING, SEND APPLICATIONS TO:

MATERIAL RADIATION PROTECTION SECTION
U.S. NUCLEAR REGULATORY COMMISSION, REGION IV
611 RYAN PLAZA DRIVE, SUITE 400
ARLINGTON, TX 76011-8064

UNITED STATES
NUCLEAR REGULATORY COMMISSION
REGION III
801 WARRENVILLE ROAD
LISLE, ILLINOIS 60532-4351

November 21, 1996

Paul M. Williams, D.O.
Radiation Safety Officer
Grim-Smith Hospital & Clinic, Inc.
d/b/a Northeast Regional Medical Cente
800 West Jefferson
Kirksville, MO 63501

SUBJECT: ACKNOWLEDGEMENT OF CORRESPONDENCE
(Letter Dated 11/15/96)

Dear Licensee:

In response to your request, we have completed the initial processing, which is an administrative review of your application for a(n):

___ New License _X_ Amendment ___ Renewal
_X_ Termination ___ Auth User (Amendment not required)
___ Other_____

No administrative deficiencies were identified during this initial review. However, it should be noted that a technical review may identify omissions in the submitted information.

It appears that your request is routine (see 1-3 below, as applicable).

1. New and amendment actions are normally processed within 90 days, unless we find major deficiencies, or policy issues requiring central program office assistance.

2. Renewal actions are normally processed within 180 days, however, under timely filing (before expiration), you may continue to operate under your existing license.

3. Termination actions are normally processed within 90 days, unless confirmatory surveys following decontamination/decommissioning activities are involved.

A copy of your correspondence has been forwarded to our Licensing Fee and Debt Collection Branch (301/415-6097) for approval of the fee category and amount, if required.

If you have a compelling safety or business-related reason for requesting expedited review, please contact the Materials Licensing Branch at (630) 829-9887. We will try to complete your request as soon as practicable. Any correspondence about this request should reference the control number.

Nuclear Materials Support Branch

Mail Control Nos.	302061	302063
License Nos.	24-05245-01	24-26702-01
	Amendment	Termination