

DEPARTMENT OF THE ARMY  
OFFICE OF THE SURGEON GENERAL  
WASHINGTON, D.C. 20310-2300

J. POTTER  
KIS

DASG-PSP-E

30 May 1985

05 AUG 5 10:32

US Nuclear Regulatory Commission  
Region II  
101 Marietta Street NW  
Suite 3100  
Atlanta, GA 30303

Gentlemen:

Two copies of a request for amendment/renewal of USNRC License No. 39-14873-01,  
Moncrief Army Community Hospital are enclosed.

Recommend approval.

Sincerely,

/s/

Enclosure

RICHARD W. FIELD  
Colonel, MSC  
Radiological Hygiene  
Consultant

8509120392 850830  
REG2 LIC30  
39-14873-01 PDR

50678 8/5/81  
Rec'd  
FEE EXEMPT

Official Copy



DEPARTMENT OF THE ARMY  
HEADQUARTERS UNITED STATES ARMY MEDICAL DEPARTMENT ACTIVITY  
FORT JACKSON, SOUTH CAROLINA 29207

0756  
9

HSXL-PRP

REPLY TO  
ATTENTION OF

9 May 1985

SUBJECT: Amendment Request for NRC License Number 39-14873-01

THRU:

Commander  
US Army Health Services Command  
ATTN: HSCL-P  
Fort Sam Houston, TX 78234-6000

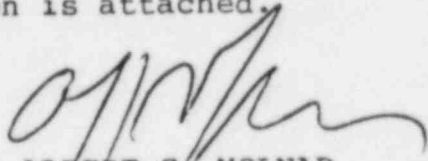
HQDA  
DASG-PSP-E  
WASH DC 20310-2300

TO:

Material Licensing Branch  
Division of Fuel Cycle and Material Safety  
Office of Nuclear Material and Safeguards  
US Nuclear Regulatory Commission  
Wash DC 20555

Request current license be amended to reflect changes in equipment and facilities. Amendment application is attached.

Atch  
as

  
ALBERT C. MOLNAR  
Colonel, Medical Corps  
Acting Commander

~~8507170568~~ 1P

NRC FORM 313M (9-81) 10 CFR 35	U.S. NUCLEAR REGULATORY COMMISSION <b>APPLICATION FOR MATERIALS LICENSE — MEDICAL</b>	Approved by OMB 3150-0041 Expires 9-30-83			
<b>INSTRUCTIONS</b> — Complete Items 1 through 26 if this is an initial application or an application for renewal of a license. Use supplemental sheets where necessary. Item 26 must be completed on all applications and signed. Retain one copy. Submit original and one copy of entire application to: Director, Office of Nuclear Materials Safety and Safeguards, U.S. Nuclear Regulatory Commission, Washington, D.C. 20555. Upon approval of this application, the applicant will receive a Materials License. An NRC Materials License is issued in accordance with the general requirements contained in Title 10, Code of Federal Regulations, Part 30, and the Licensee is subject to Title 10, Code of Federal Regulations, Parts 19, 20 and 35 and the license fee provision of Title 10, Code of Federal Regulations, Part 170. The license fee category should be stated in Item 26 and the appropriate fee enclosed.					
1.a. NAME AND MAILING ADDRESS OF APPLICANT (institution, firm, clinic, physician, etc.) INCLUDE ZIP CODE  Department of the Army Nuclear Medicine Service Moncrief Army Community Hospital Fort Jackson, SC 29207-5720 TELEPHONE NO.: AREA CODE 803 751 2383		1.b. STREET ADDRESS(ES) AT WHICH RADIOACTIVE MATERIAL WILL BE USED (If different from 1.a.) INCLUDE ZIP CODE  Same as 1a.			
2. PERSON TO CONTACT REGARDING THIS APPLICATION  CPT J. R. CARTER  TELEPHONE NO.: AREA CODE (803) 751 4552/2267		3. THIS IS AN APPLICATION FOR: (Check appropriate item) a. <input type="checkbox"/> NEW LICENSE b. <input checked="" type="checkbox"/> AMENDMENT TO LICENSE NO. 39-14873-01 c. <input type="checkbox"/> RENEWAL OF LICENSE NO.			
4. INDIVIDUAL USERS (Name individuals who will use or directly supervise use of radioactive material. Complete Supplements A and B for each individual.)  NO CHANGE		5. RADIATION SAFETY OFFICER (RSO) (Name of person designated as radiation safety officer. If other than individual user, complete resume of training and experience as in Supplement A.)  NO CHANGE			
6.a. RADIOACTIVE MATERIAL FOR MEDICAL USE <span style="float: right;">NO CHANGE</span>					
RADIOACTIVE MATERIAL LISTED IN:	ITEMS DESIRED "X"	MAXIMUM POSSESSION LIMITS (In millicuries)	ADDITIONAL ITEMS:	MARK ITEMS DESIRED "X"	MAXIMUM POSSESSION LIMITS (In millicuries)
10 CFR 31.11 FOR IN VITRO STUDIES			IODINE-131 AS IODIDE FOR TREATMENT OF HYPERTHYROIDISM		
10 CFR 35.100, SCHEDULE A, GROUP I		AS NEEDED	PHOSPHORUS-32 AS SOLUBLE PHOSPHATE FOR TREATMENT OF POLYCYTHEMIA VERA, LEUKEMIA AND BONE METASTASES		
10 CFR 35.100, SCHEDULE A, GROUP II		AS NEEDED	PHOSPHORUS-32 AS COLLOIDAL CHROMIC PHOSPHATE FOR INTRACAVITARY TREATMENT OF MALIGNANT EFFUSIONS.		
10 CFR 35.100, SCHEDULE A, GROUP III			GOLD-198 AS COLLOID FOR INTRACAVITARY TREATMENT OF MALIGNANT EFFUSIONS.		
10 CFR 35.100, SCHEDULE A, GROUP IV		AS NEEDED	IODINE-131 AS IODIDE FOR TREATMENT OF THYROID CARCINOMA		
10 CFR 35.100, SCHEDULE A, GROUP V		AS NEEDED	XENON-133 AS GAS OR GAS IN SALINE FOR BLOOD FLOW STUDIES AND PULMONARY FUNCTION STUDIES.		
10 CFR 35.100, SCHEDULE A, GROUP VI					
6.b. RADIOACTIVE MATERIAL FOR USES NOT LISTED IN ITEM 6.a. (Sealed sources up to 3 mCi used for calibration and reference standards are authorized under Section 35.14(d), 10 CFR Part 35, and NEED NOT BE LISTED.)					
ELEMENT AND MASS NUMBER	CHEMICAL AND/OR PHYSICAL FORM	MAXIMUM NUMBER OF MILLICURIES OF EACH FORM	DESCRIBE PURPOSE OF USE		

# INFORMATION REQUIRED FOR ITEMS 7 THROUGH 23

For Items 7 through 23, check the appropriate box(es) and submit a detailed description of all the requested information. Begin each item on a separate sheet. Identify the item number and the date of the application in the lower right corner of each page. If you indicate that an appendix to the medical licensing guide will be followed, do not submit the pages, but specify the revision number and date of the referenced guide: Regulatory Guide 10.8 Rev. 1 Date: Oct 1980

7. MEDICAL ISOTOPES COMMITTEE <u>NO CHANGE</u>		15. GENERAL RULES FOR THE SAFE USE OF RADIOACTIVE MATERIAL <i>(Check One)</i> <u>NO CHANGE</u>	
<input type="checkbox"/>	Names and Specialties Attached; and	<input type="checkbox"/>	Appendix G Rules Followed; or
<input type="checkbox"/>	Duties as in Appendix B; or _____ <i>(Check One)</i>	<input type="checkbox"/>	Equivalent Rules Attached
<input type="checkbox"/>	Equivalent Duties Attached	16. EMERGENCY PROCEDURES <i>(Check One)</i> <u>NO CHANGE</u>	
8. TRAINING AND EXPERIENCE <u>NO CHANGE</u>		<input type="checkbox"/>	Appendix H Procedures Followed; or
<input type="checkbox"/>	Supplements A & B Attached for Each Individual User; and	<input type="checkbox"/>	Equivalent Procedures Attached
<input type="checkbox"/>	Supplement A Attached for RSO.	17. AREA SURVEY PROCEDURES <i>(Check One)</i> <u>NO CHANGE</u>	
9. INSTRUMENTATION <i>(Check One)</i> <u>NO CHANGE</u>		<input type="checkbox"/>	Appendix I Procedures Followed; or
<input type="checkbox"/>	Appendix C Form Attached; or	<input type="checkbox"/>	Equivalent Procedures Attached
<input type="checkbox"/>	List by Name and Model Number	18. WASTE DISPOSAL <i>(Check One)</i> <u>NO CHANGE</u>	
10. CALIBRATION OF INSTRUMENTS <u>NO CHANGE</u>		<input type="checkbox"/>	Appendix J Form Attached; or
<input type="checkbox"/>	Appendix D Procedures Followed for Survey Instruments; or _____ <i>(Check One)</i>	<input type="checkbox"/>	Equivalent Information Attached
<input type="checkbox"/>	Equivalent Procedures Attached; and	19. THERAPEUTIC USE OF RADIOPHARMACEUTICALS <i>(Check One)</i> <u>NO CHANGE</u>	
<input type="checkbox"/>	Appendix D Procedures Followed for Dose Calibrator; or _____ <i>(Check One)</i>	<input type="checkbox"/>	Appendix K Procedures Followed; or
<input type="checkbox"/>	Equivalent Procedures Attached	<input type="checkbox"/>	Equivalent Procedures Attached
11. FACILITIES AND EQUIPMENT <u>TAB A</u>		20. THERAPEUTIC USE OF SEALED SOURCES <u>NO CHANGE</u>	
<input type="checkbox"/>	Description and Diagram Attached	<input type="checkbox"/>	Detailed Information Attached; and
12. PERSONNEL TRAINING PROGRAM <u>NO CHANGE</u>		<input type="checkbox"/>	Appendix L Procedures Followed; or _____ <i>(Check One)</i>
<input type="checkbox"/>	Description of Training Attached	<input type="checkbox"/>	Equivalent Procedures Attached
13. PROCEDURES FOR ORDERING AND RECEIVING RADIOACTIVE MATERIAL <u>NO CHANGE</u>		21. PROCEDURES AND PRECAUTIONS FOR USE OF RADIOACTIVE GASES (e.g., Xenon - 133) <u>NO CHANGE</u>	
<input type="checkbox"/>	Detailed Information Attached	<input type="checkbox"/>	Detailed Information Attached
14. PROCEDURES FOR SAFELY OPENING PACKAGES CONTAINING RADIOACTIVE MATERIALS <i>(Check One)</i> <u>NO CHANGE</u>		22. PROCEDURES AND PRECAUTIONS FOR USE OF RADIOACTIVE MATERIAL IN ANIMALS <u>NO CHANGE</u>	
<input type="checkbox"/>	Appendix F Procedures Followed; or	<input type="checkbox"/>	Detailed Information Attached
<input type="checkbox"/>	Equivalent Procedures Attached	23. PROCEDURES AND PRECAUTIONS FOR USE OF RADIOACTIVE MATERIAL SPECIFIED IN ITEM 6.b <u>NO CHANGE</u>	
<input type="checkbox"/>		<input type="checkbox"/>	Detailed Information Attached

## 24. PERSONNEL MONITORING DEVICES

NO CHANGE

TYPE (Check appropriate box)		SUPPLIER	EXCHANGE FREQUENCY
a. WHOLE BODY	FILM		
	TLD		
	OTHER (Specify)		
b. FINGER	FILM		
	TLD		
	OTHER (Specify)		
c. WRIST	FILM		
	TLD		
	OTHER (Specify)		

d. OTHER (Specify)

## 25. FOR PRIVATE PRACTICE APPLICANTS ONLY

a. HOSPITAL AGREEING TO ACCEPT PATIENTS CONTAINING RADIOACTIVE MATERIAL

NAME OF HOSPITAL

b. ATTACH A COPY OF THE AGREEMENT LETTER  
SIGNED BY THE HOSPITAL ADMINISTRATOR.

MAILING ADDRESS

c. WHEN REQUESTING THERAPY PROCEDURES,  
ATTACH A COPY OF RADIATION SAFETY PRECAU-  
TIONS TO BE TAKEN AND LIST AVAILABLE  
RADIATION DETECTION INSTRUMENTS.

CITY

STATE

ZIP CODE

## 26. CERTIFICATE

(This item must be completed by applicant)

The applicant and any official executing this certificate on behalf of the applicant named in Item 1a certify that this application is prepared in conformity with Title 10, Code of Federal Regulations, Parts 30 and 35, and that all information contained herein, including any supplements attached hereto, is true and correct to the best of our knowledge and belief.

a. LICENSE FEE REQUIRED  
(See Section 170.31, 10 CFR 170)

b. APPLICANT OR CERTIFYING OFFICIAL (Signature)

(1) NAME (Type of Print)

ALBERT C. MOLNAR, Colonel, MC

(2) TITLE

Acting Commander

(1) LICENSE FEE CATEGORY:

EXEMPT

c. DATE

9 May 1985

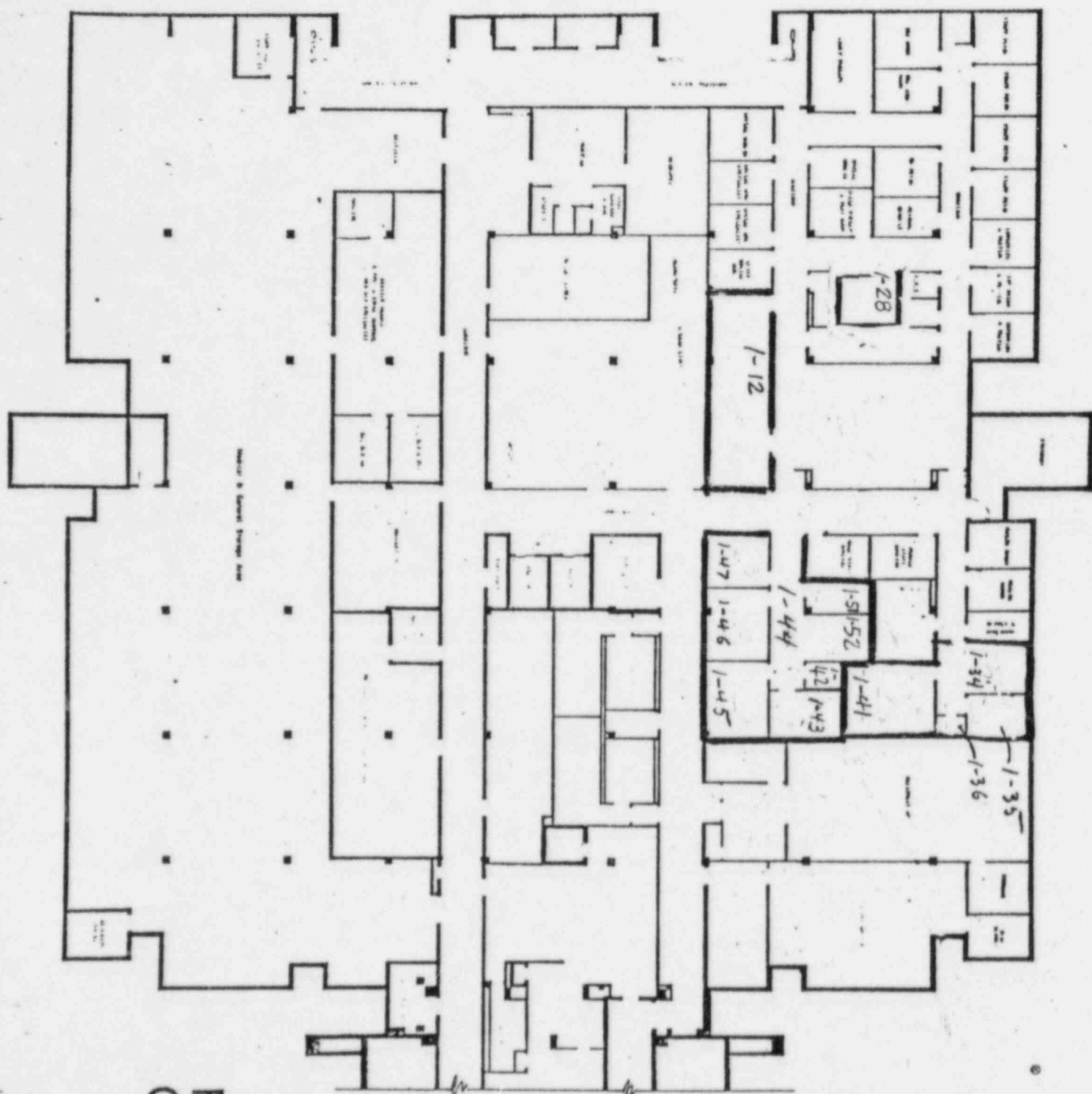
(2) LICENSE FEE ENCLOSED: \$

TAB A

1. Request approval of new facility site and layout for the MACH Nuclear Medicine Svc, In Vitro Laboratory as depicted in the attached diagrams.
2. The proposed move of the In Vitro Lab and administrative offices will consolidate the Nuclear Medicine Svc in one clinical location preventing the routine transportation of radiopharmaceuticals across public corridors. This move should reduce the risk of accidental contamination due to material movements within the hospital.
3. In accordance with US NRC Reg Guide 10.8 Appendix N para 1j(3) a close out survey of the In Vitro lab site will be conducted insuring no radioactive sources remain and there is no removable contamination present prior to release for nonrestricted use. Until the close-out survey is completed documenting that all sources have been removed and that the lab area is free of contamination this area will remain a restricted area and will be under the supervision of the Radiation Protection Officer.
4. Upon completion of the close out survey a copy of the survey will be forwarded to your office if requested.



# EXISTING FACILITY

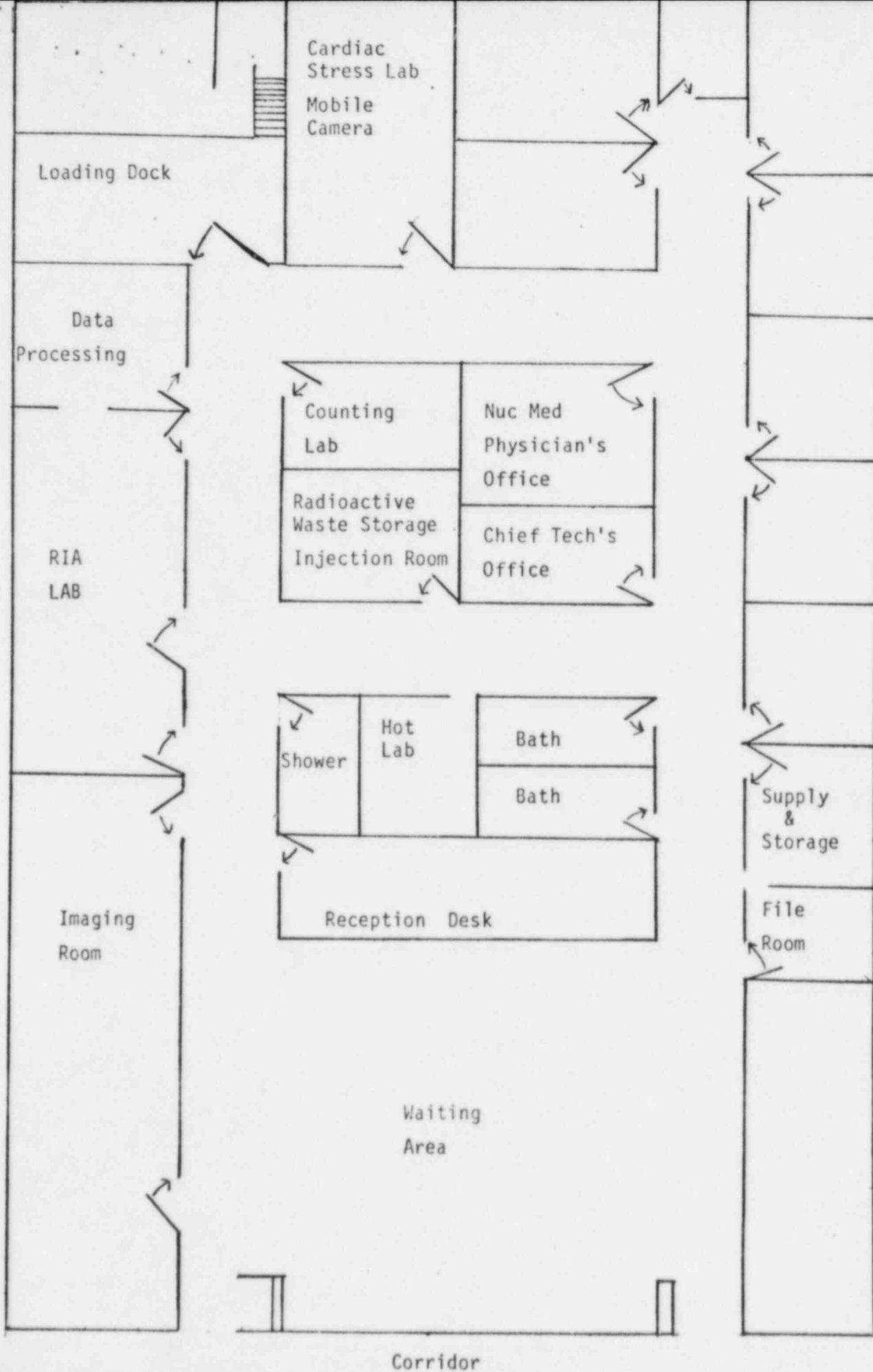


## FIRST FLOOR CLINIC WING

"Rm 1-12--Imaging -  
1-28--Radiopharmach/  
Hot Lab  
1-47--Drs. Office  
1-46--Reception  
1-45--Imaging -  
1-44--Hallway  
1-43--Data Processing

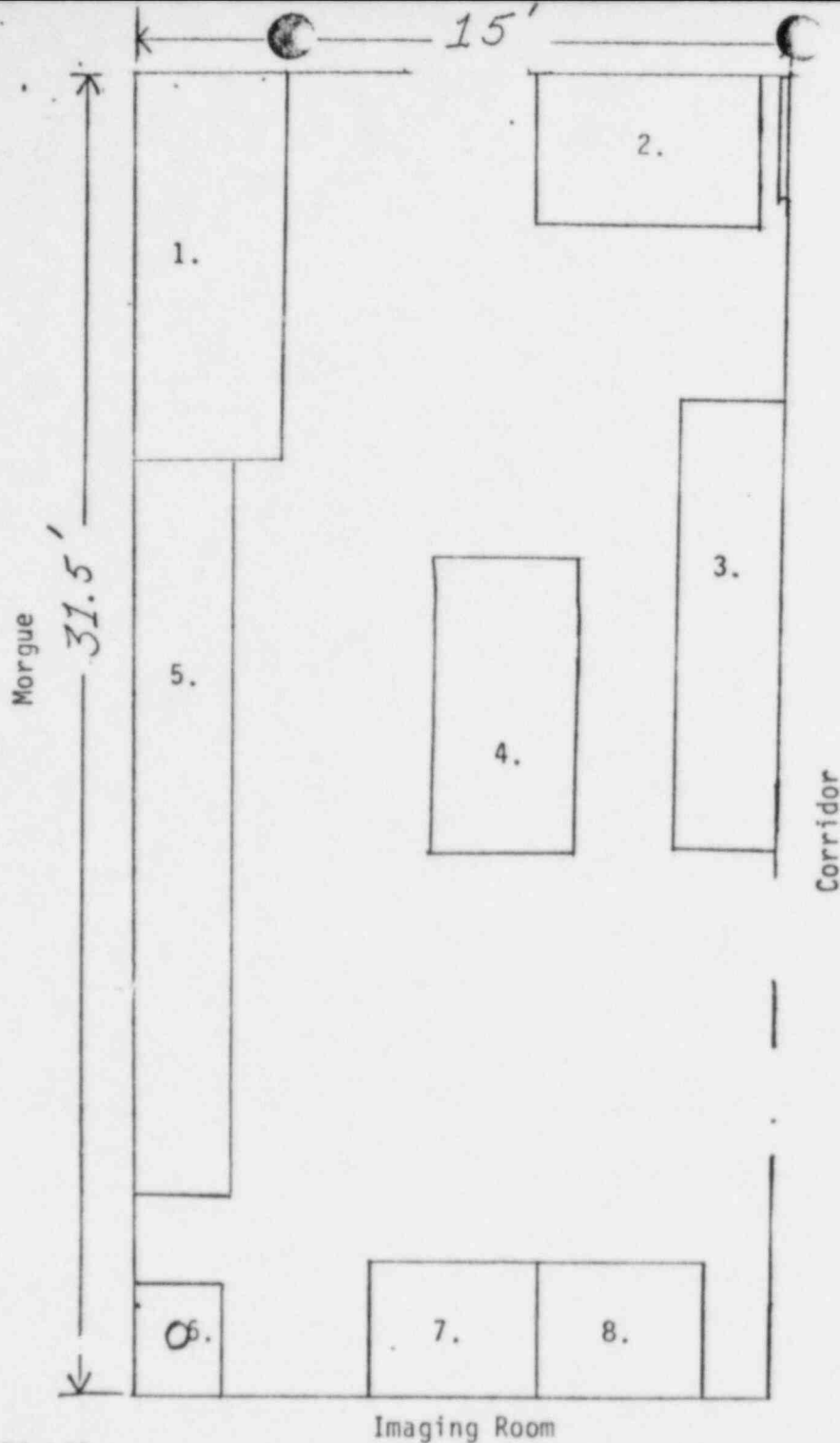
SEE 'FIRST FLOOR TOWER' FOR CONTINUATION

Rm 1-51--Injection Room -  
1-52--Pt. waiting  
1-42--Toilet(Staff)  
1-41--In-vitro Lab -  
1-34--Counting Lab -  
1-35--Office, Chief Tech.  
1-36--Closet



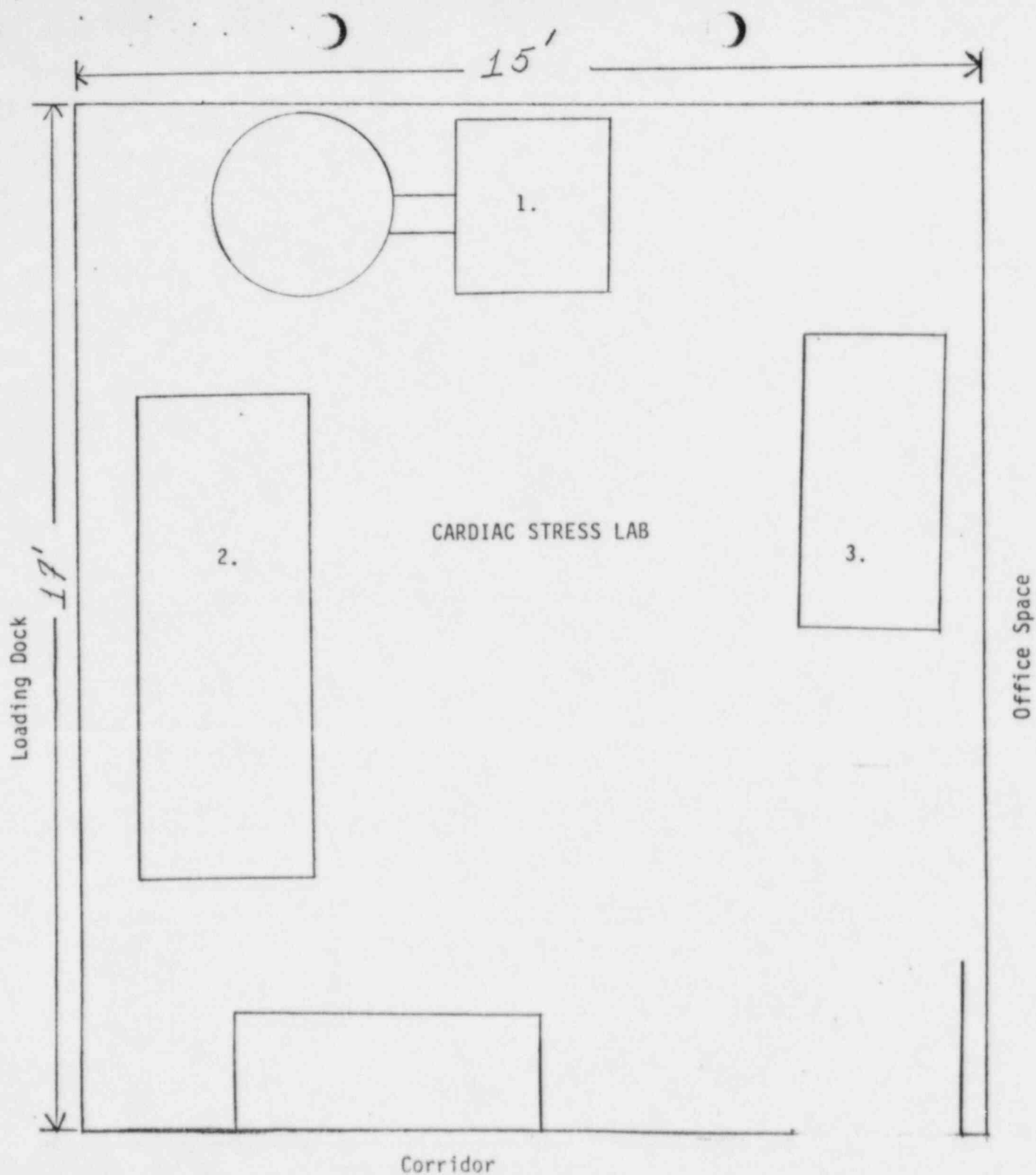
PROPOSED LAYOUT



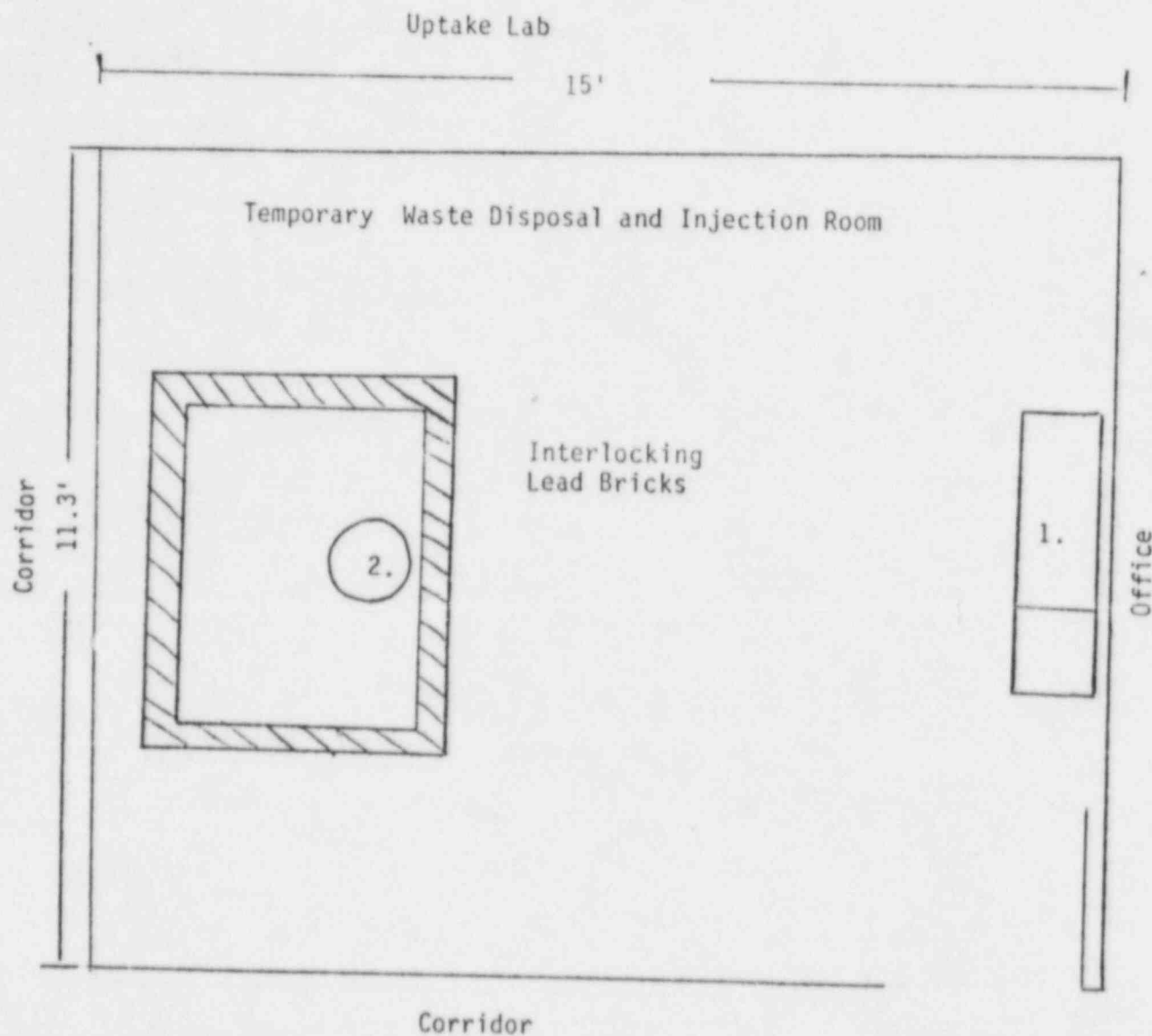


# IN VITRO LAB

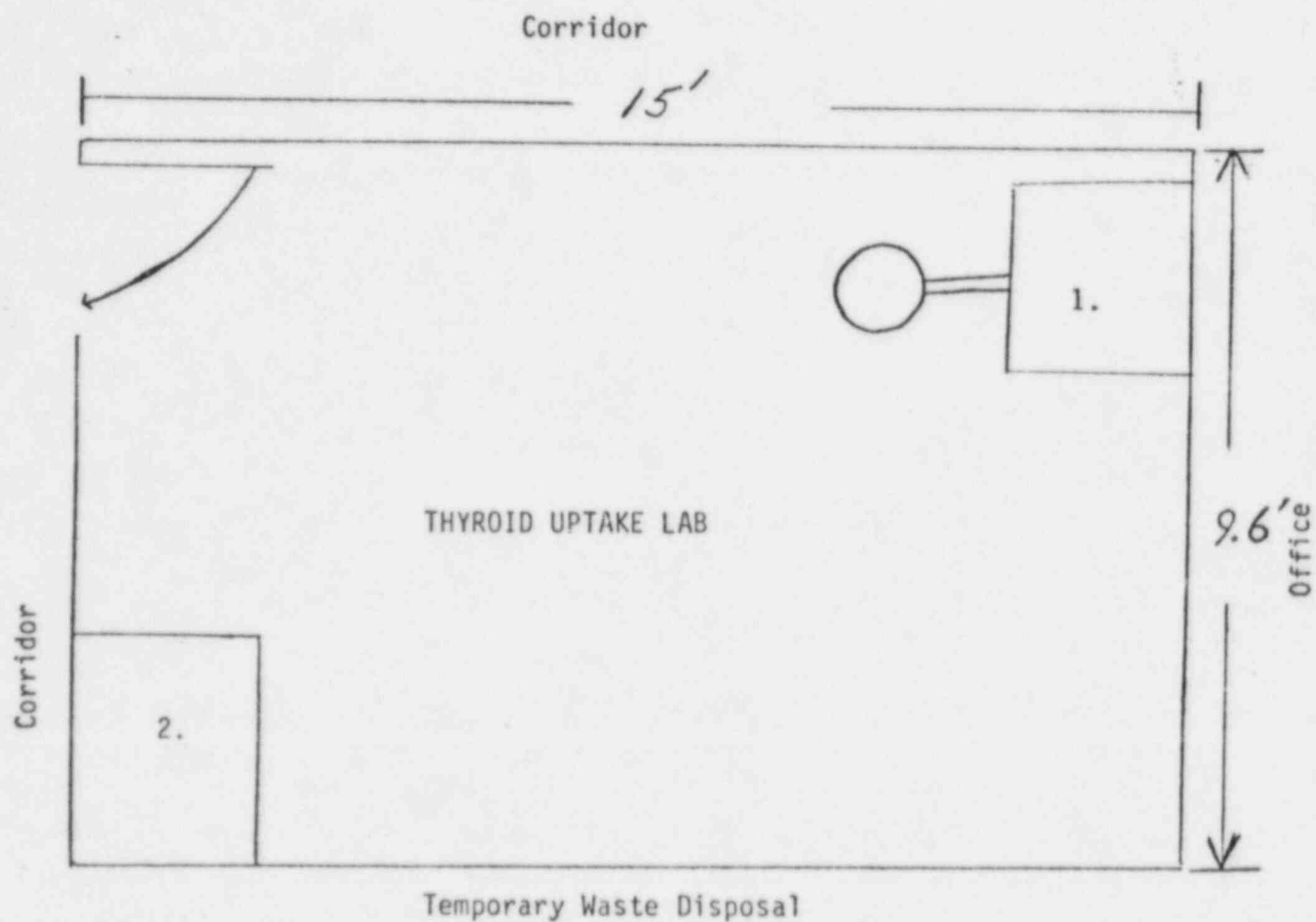
1. Concept 4.
2. 4/600 Gamma Counter.
3. Counter Top.
4. Couter Top.
5. Counter Top.
6. Hot Sink.
7. Freezer.
8. Refrigerator.



1. Portable Gamma Camera.
2. Treadmill.
3. Exercise Bicycle.



1. Injection Chair.
2. Technecium Generator and Generator Housing. (Generator has .25 Pb equivalent and Generator Housing has .25 Pb equivalent)



- 1. Uptake Probe.
- 2. Centrifuge.